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Les documents fixés à cette attestation sont conformes à la version initialement déposée de la demande de brevet européen spécifiée à la page suivante.

Patentanmeldung Nr. Patent application No. Demande de brevet n°

02290822.2

**PRIORITY
DOCUMENT**

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For the President of the European Patent Office

Le Président de l'Office européen des brevets
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R C van Dijk

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151 85 Södertälje
SUEDE

Bezeichnung der Erfindung/Title of the invention/Titre de l'invention:
(Falls die Bezeichnung der Erfindung nicht angegeben ist, siehe Beschreibung.
If no title is shown please refer to the description.
Si aucun titre n'est indiqué se referer à la description.)

Chemical compounds

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AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE TR

- 1 -

CHEMICAL COMPOUNDS

This invention relates to vascular damaging agents and their uses. In particular it relates to certain compounds which may be of use as vascular damaging agents, to methods for preparing the compounds, to their use as medicaments (including methods for the treatment of angiogenesis or disease states associated with angiogenesis) and to pharmaceutical compositions containing them. The invention also relates to the use of such compounds in the manufacture of medicaments for the production of anti-angiogenic and/or anti-vascular effects.

Normal angiogenesis plays an important role in a variety of processes including embryonic development, wound healing and several components of female reproductive function. Undesirable or pathological angiogenesis has been associated with disease states including diabetic retinopathy, psoriasis, cancer, rheumatoid arthritis, atheroma, Kaposi's sarcoma and haemangioma (Fan et al, 1995, Trends Pharmacol. Sci. 16: 57-66; Folkman, 1995, Nature Medicine 1: 27-31). Formation of new vasculature by angiogenesis is a key pathological feature of several diseases (J. Folkman, New England Journal of Medicine 333, 1757-1763 (1995)). For example, for a solid tumour to grow it must develop its own blood supply upon which it depends critically for the provision of oxygen and nutrients; if this blood supply is mechanically shut off the tumour undergoes necrotic death. Neovascularisation is also a clinical feature of skin lesions in psoriasis, of the invasive pannus in the joints of rheumatoid arthritis patients and of atherosclerotic plaques. Retinal neovascularisation is pathological in macular degeneration and in diabetic retinopathy.

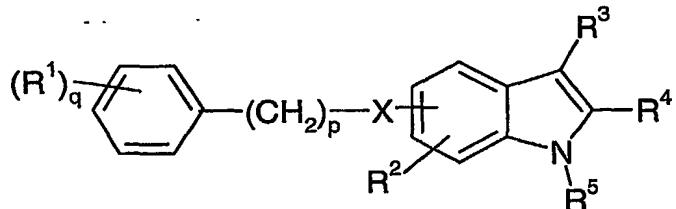
Reversal of neovascularisation by damaging the newly-formed vascular endothelium is therefore expected to have a beneficial therapeutic effect. Such vascular-damaging activity would clearly be of value in the treatment of disease states associated with angiogenesis such as cancer, diabetes, psoriasis, rheumatoid arthritis, Kaposi's sarcoma, haemangioma, acute and chronic nephropathies, atheroma, arterial restenosis, autoimmune diseases, acute inflammation, endometriosis, dysfunctional uterine bleeding and ocular diseases with retinal vessel proliferation.

Certain known compounds that cause selective destruction of tumour vasculature have been reported, *in vitro* and at non-cytotoxic concentrations, to cause effects on proliferating endothelial cells, ie, cell detachment [Blakey D C et al, *Proceedings of the American*

*Association for Cancer Research, 41, 329, 2000 abstract 2086] and changes in cell shape [Davis P D et al, *Proceedings of the American Association for Cancer Research, 41, 329, 2000 abstract 2085; Chaplin D J & Dougherty G J, Br J Cancer, 80, Suppl 1, 57-64, 1999*. It can therefore be expected that these compounds will have damaging effects on newly-formed 5 vasculature, for example the vasculature of tumours. It can reasonably be predicted, for example, that they will be capable of causing selective destruction of tumour vasculature, both *in vitro* and *in vivo*. Destruction of tumour vasculature in turn leads to a reduction in tumour blood flow and to tumour cell death due to starvation of oxygen and nutrients, ie, to anti-tumour activity [Davis P D et al; Chaplin D J & Dougherty G J; Blakey D C et al, all *supra*].*

10 Compounds with this activity have also been described in International Patent Application WO 99/02166 (Angiogene Pharmaceuticals), International Patent Application WO00/40529 (Angiogene Pharmaceuticals) and International Patent Application WO 00/41669 (Angiogene Pharmaceuticals).

We have identified a class of indole compounds with vascular damaging activity.
 15 Thus, according to the first feature of the present invention there is provided the use of a compound of Formula (I), for the manufacture of a medicament to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis



Formula (I)

20

wherein:

R¹ is independently selected from halo, hydroxy, amino, alkanoylamino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;

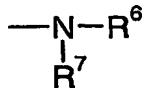
25 X is selected from: —O—, —S—, —SO— or —SO₂—;

R² is selected from: hydrogen, C₁₋₄alkyl or C₁₋₄alkoxy;

R³ and R⁴ are independently selected from: hydrogen, C₁₋₄alkyl, C₁₋₄alkanoyl,

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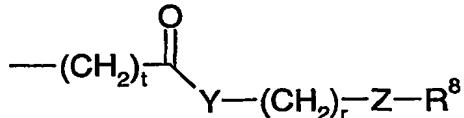
aminoC₁₋₄alkyl, carbamoyl, carbamoylC₁₋₄alkyl, cyano, cyanoC₁₋₄alkyl, hydroxy, hydroxyC₁₋₄alkyl, or a group of Formula (II):



Formula (II)

5 R⁶ is hydrogen or C₁₋₄alkyl;

R⁵ and R⁷ are independently selected from hydrogen, C₁₋₄alkyl or a group of Formula (III):



Formula (III)

wherein Y is selected from —NH—, —O— or a bond;

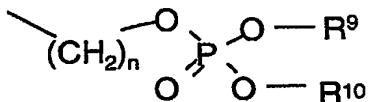
10 Z is selected from —NH—, —O—, —C(O)— or a bond;

r is an integer from 0 to 4;

t is an integer from 0 to 1;

R⁸ is hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy, aryl, 5 or 6 membered heterocyclyl, 5- or 6-membered heteroaryl, wherein aryl, heteroaryl or heterocyclyl are optionally substituted by C₁₋₄alkyl, C₁₋₄alkoxy, or a group of

15 Formula (IV):



Formula (IV)

wherein n is an integer from 1 to 6, and;

20 R⁹ and R¹⁰ are independently selected from hydrogen, C₁₋₄alkyl or aryl; and

p is an integer between 0 and 1; and

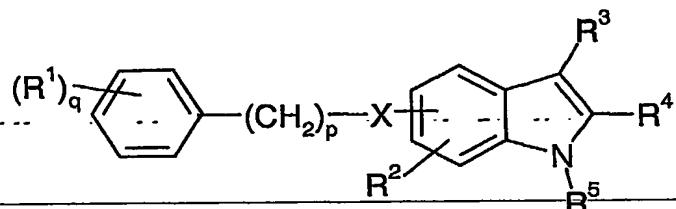
q is an integer between 0 and 3;

with the proviso that:

25 (i) when R³ is cyano then R⁴ cannot be a group of Formula (II), and

(ii) when q is 0, R³ is cyano and X is —S— then R⁴ is other than amino; or a salt, pro-drug or solvate thereof.

According to a further aspect of the first feature of the invention there is provided the use of a compound of Formula (Ia), for the manufacture of a medicament to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis:



5

Formula (Ia)

wherein:

R^1 is independently selected from hydroxy, amino, alkanoylamino, — OPO_3H_2 , or C_{1-4} alkoxy,

wherein the amino group is optionally substituted by an amino acid residue and the

10 hydroxy group is optionally esterified;

wherein: X , p , q , R^2 , R^3 , R^4 , and R^5 are as defined for Formula (I) above;

with the proviso that:

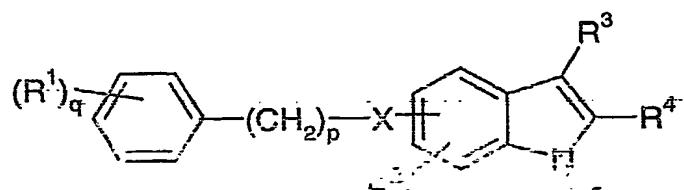
(i) when R^3 is cyano then R^4 cannot be a group of Formula (II), and

or a salt, pro-drug or solvate thereof.

15 According to a further aspect of the first feature of the invention there is provided a method of treatment, in a warm-blooded animal, to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis comprising administering to said warm-blooded animal a therapeutically (including prophylactically) effective amount of a compound of Formula (I) or Formula (Ia), or a pharmaceutically acceptable salt, pro-drug or solvate thereof.

20 Preferably a warm-blooded animal is a human.

A further feature of the invention provides a group of indole compounds for use in medicine. Thus, according to a second feature of the present invention there is provided the use of a compound of Formula (V) as a medicament, wherein:



- 5 -

Formula (V)

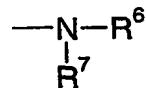
R^1 is independently selected from halo, hydroxy, amino, alkanoylamino, $-\text{OPO}_3\text{H}_2$, or $C_{1-4}\text{alkoxy}$, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;

5 X is selected from: $-\text{O}-$, $-\text{S}-$, $-\text{SO}-$ or $-\text{SO}_2-$;

R^2 is selected from: hydrogen, $C_{1-4}\text{alkyl}$ or $C_{1-4}\text{alkoxy}$;

R^3 and R^4 are independently selected from: hydrogen, $C_{1-4}\text{alkyl}$, $C_{1-4}\text{alkanoyl}$, $C_{1-4}\text{alkoxycarbonyl}$, $C_{1-4}\text{alkoxycarbonyl}C_{1-4}\text{alkyl}$, $C_{1-4}\text{alkoxycarbonylamino}$, amino, amino $C_{1-4}\text{alkyl}$, carbamoyl, carbamoyl $C_{1-4}\text{alkyl}$, cyano, cyano $C_{1-4}\text{alkyl}$, hydroxy,

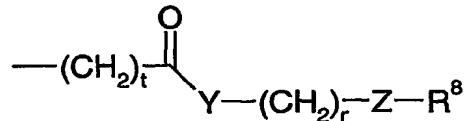
10 hydroxy $C_{1-4}\text{alkyl}$; or a group of Formula (II):



Formula (II)

R^6 is hydrogen or $C_{1-4}\text{alkyl}$;

R^5 and R^7 are independently selected from hydrogen, $C_{1-4}\text{alkyl}$ or a group of Formula (III):



15

Formula (III)

wherein Y is selected from $-\text{NH}-$, $-\text{O}-$ or a bond;

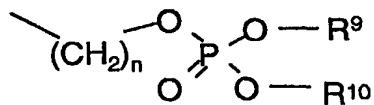
Z is selected from $-\text{NH}-$, $-\text{O}-$, $-\text{C(O)}-$ or a bond;

r is an integer from 0 to 4;

20

t is an integer from 0 to 1;

R^8 is hydrogen, $C_{1-4}\text{alkyl}$, $C_{1-4}\text{alkoxy}$, aryl, 5 or 6 membered heterocyclyl, 5- or 6-membered heteroaryl, wherein aryl, heteroaryl or heterocyclyl is optionally substituted by $C_{1-4}\text{alkyl}$ or $C_{1-4}\text{alkoxy}$, or a group of Formula (IV):



25

Formula (IV)

wherein n is an integer of from 1 to 6, and;

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R^9 and R^{10} are independently selected from hydrogen or C_{1-4} alkyl or aryl; and

p is an integer between 0 and 1; and

q is an integer between 0 and 3;

5 with the proviso that

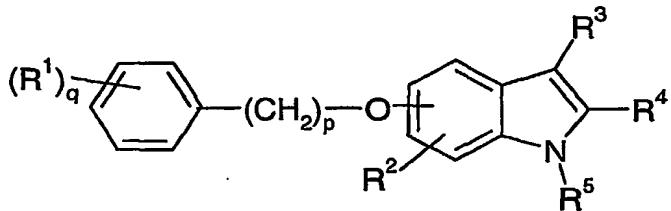
- (i) when R^3 is cyano then R^4 cannot be a group of Formula (II);
- (ii) when q is 0 and X is —S— then R^3 is other than cyano;
- (iii) when q is 0, p is 1, R^2 is hydrogen or 5-methoxy, R^3 is cyanomethyl and R^4 is hydrogen or R^3 is hydrogen and R^4 is cyanomethyl then R^5 cannot be hydrogen methyl or acetyl;
- (iv) when $(R^1)_q$ is 4-methoxy, 4-amino or 3,4,5-trimethoxy, p is 0 or 1, R^2 is hydrogen, 4-chloro or 5-methoxy, R^3 is hydrogen, cyanomethyl or 2-aminoethyl, R^4 is hydrogen or ethoxycarbonyl then R^5 cannot be hydrogen or methyl;
- (v) when q is 0, p is 0 or 1, R^2 is hydrogen, 7-methyl, 5-methoxy or 6-methoxy, R^3 is aminomethyl, 2-aminoethyl, 2-aminopropyl or 2-aminobutyl, R^4 is hydrogen or methyl, then R^5 cannot be hydrogen, methyl, n-butyl or acetyl;
- (vi) when q is 0, p is 0 or 1, R^2 is hydrogen, R^3 is methyl, ethyl, hydroxy, hydroxymethyl, 2-hydroxyethyl or 1-methylethoxy, R^4 is hydrogen, methyl, ethoxycarbonyl or *tert*-butoxycarbonyl or carbamoyl, then R^5 cannot be hydrogen, methyl or acetyl;
- (vii) when q is 0, p is 0 or 1, R^2 is hydrogen or 5-methoxy, R^3 is hydrogen or bromo, R^4 is hydrogen, methyl, methoxycarbonyl, ethoxycarbonyl, *tert*-butoxycarbonyl or hydroxymethyl then R^5 cannot be methyl or n-butyl;
- 25 and
- (viii) when q is 0, p is 1, R^2 is hydrogen, 6-methyl, 7-methyl, or 5-methoxy, R^3 is hydrogen, R^4 is hydrogen, methoxycarbonyl, ethoxycarbonyl, or 2-aminoethyl, then R^5 cannot be hydrogen;

or a salt, pro-drug or solvate thereof;

30 According to a further aspect of the second feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (V) or pharmaceutically-acceptable salt, pro-drug or solvate thereof.

According to a further aspect of the second feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (V), or a pharmaceutically-acceptable salt, pro-drug or solvate thereof, in admixture with a pharmaceutically-acceptable diluent or carrier.

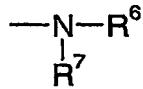
- 5 The invention also provides a novel group of indole compounds. Thus, according to a third feature of the invention there is provided a compound of Formula (VI)



Formula (VI)

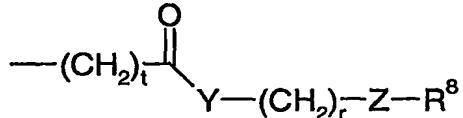
wherein:

- 10 R¹ is independently selected from halo, hydroxy, amino, alkanoylamino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;
- R² is selected from: hydrogen, C₁₋₄alkyl or C₁₋₄alkoxy;
- R³ and R⁴ are independently selected from: hydrogen, C₁₋₄alkyl, C₁₋₄alkanoyl,
- 15 C₁₋₄alkoxycarbonyl, C₁₋₄alkoxycarbonylC₁₋₄alkyl, C₁₋₄alkoxycarbonylamino, amino, aminoC₁₋₄alkyl, carbamoyl, carbamoylC₁₋₄alkyl, cyano, cyanoC₁₋₄alkyl, hydroxy, hydroxyC₁₋₄alkyl; or a group of Formula (II):



Formula (II)

- 20 R⁶ is hydrogen or C₁₋₄alkyl;
- R⁵ and R⁷ are independently selected from hydrogen, C₁₋₄alkyl or a group of Formula (III):



Formula (III)

wherein Y is selected from —NH—, —O— or a bond;

- 25 Z is selected from —NH—, —O—, —C(O)— or a bond;

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r is an integer from 0 to 4;

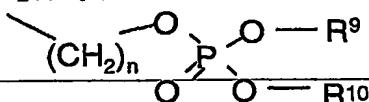
t is an integer from 0 to 1;

R⁸ is hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy, aryl, 5 or 6 membered heterocyclyl,

5 5- or 6-membered heteroaryl, wherein aryl, heteroaryl or heterocyclyl

is optionally substituted by C₁₋₄alkyl or C₁₋₄alkoxy, or a group of

Formula (IV):



Formula (IV)

wherein n is an integer of from 1 to 6, and;

10 R⁹ and R¹⁰ are independently selected from hydrogen or

C₁₋₄alkyl or aryl; and

p is an integer between 0 and 1; and

q is an integer between 0 and 3;

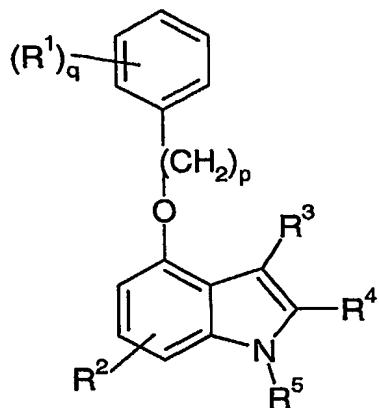
with the proviso that

- 15 (i) when R³ is cyano then R⁴ cannot be a group of Formula (II);
- (ii) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is cyanomethyl and R⁴ is hydrogen or R³ is hydrogen and R⁴ is cyanomethyl then R⁵ cannot be hydrogen methyl or acetyl;
- (iii) when (R¹)_q is 4-methoxy, 4-amino, or 3,4,5-trimethoxy, p is 0 or 1, R² is hydrogen, 4-chloro or 5-methoxy, R³ is hydrogen, cyanomethyl or 2-aminoethyl, R⁴ is hydrogen or ethoxycarbonyl then R⁵ cannot be hydrogen or methyl;
- 20 (iv) when q is 0, p is 0 or 1, R² is hydrogen, 7-methyl, 5-methoxy or 6-methoxy, R³ is aminomethyl, 2-aminoethyl, 2-aminopropyl or 2-aminobutyl, R⁴ is hydrogen or methyl, then R⁵ cannot be hydrogen, methyl, n-butyl or acetyl;
- (v) when q is 0, p is 1, R² is hydrogen, R³ is methyl, ethyl, hydroxy, hydroxymethyl, 2-hydroxyethyl or 1-methylethoxy, R⁴ is hydrogen, methyl, ethoxycarbonyl or tert-butoxycarbonyl or carbamoyl, then R⁵ cannot be hydrogen, methyl or acetyl;
- 25 (vi) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is hydrogen or bromo, R⁴ is hydrogen, methyl, methoxycarbonyl, ethoxycarbonyl, tert-butoxycarbonyl or hydroxymethyl then R⁵ cannot be methyl or n-butyl; and

- (vii) when q is 0, p is 1, R² is hydrogen, 6-methyl, 7-methyl, or 5-methoxy, R³ is hydrogen, R⁴ is methoxycarbonyl, ethoxycarbonyl, or 2-aminoethyl, then R⁵ cannot be hydrogen;

or a salt, pro-drug or solvate thereof;

5 According to a further aspect of the third feature of the invention there is provided a compound of Formula (VIa)



Formula (VIa)

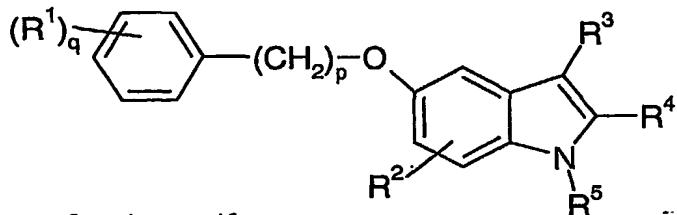
wherein: p, q, R¹, R², R³, R⁴, and R⁵ are as defined for Formula (VI) above

10 with the proviso that

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II);
- (ii) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is cyanomethyl and R⁴ is hydrogen or R³ is hydrogen and R⁴ is cyanomethyl then R⁵ cannot be hydrogen;
- (iii) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is aminomethyl, 2-aminoethyl, 2-aminopropyl, 2-aminobutyl, methoxycarbonyl or methyl, R⁴ is hydrogen or methyl, then R⁵ cannot be hydrogen, methyl or n-butyl; and
- (iv) when q is 0, p is 1, 2 is hydrogen or 5-methoxy, R³ is hydrogen, R⁴ is hydrogen, methyl, hydroxymethyl or methoxycarbonyl then R⁵ cannot be hydrogen, methyl or n-butyl.

15 20 or a salt, pro-drug or solvate thereof;

According to a further aspect of the third feature of the invention there is provided a compound of Formula (VIb)



Formula (VIb)

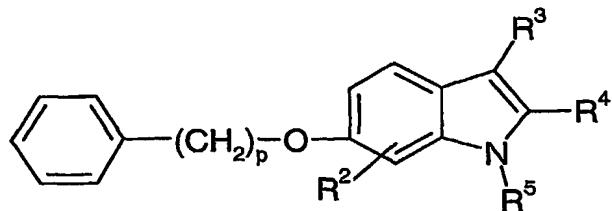
5 wherein: p, q, R¹, R², R³, R⁴, and R⁵ are as defined for Formula (VI) above

with the proviso that

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II);
- (ii) when (R¹)_q is 4-methoxy, p is 0, R² is hydrogen, R³ is hydrogen, cyanomethyl or 2-aminoethyl R⁴ is hydrogen or ethoxycarbonyl, then R⁵ cannot be hydrogen;
- 10 (iii) when q is 0, p is 1, R² is hydrogen, R³ is cyanomethyl and R⁴ is hydrogen or R³ is hydrogen and R⁴ is cyanomethyl then R⁵ cannot be hydrogen, methyl or acetyl;
- (iv) when q is 0, p is 0 or 1, R² is hydrogen, 6-methoxy or 7-methyl, R³ is 2-aminoethyl or 2-aminopropyl, R⁴ is hydrogen or methyl, then R⁵ cannot be hydrogen, methyl or acetyl;
- 15 (v) when q is 0, p is 1, R² is hydrogen, R³ is methyl, ethyl, bromo, hydroxy, hydroxymethyl, 2-hydroxyethyl, 1-methylethoxy, acetyl, ethoxycarbonyl or ethoxycarbonylethyl, R⁴ is hydrogen, methyl, carbamoyl, ethoxycarbonyl or *tert*-butoxycarbonyl, then R⁵ cannot be hydrogen or methyl; and
- (vi) when q is 0, R¹ is hydrogen, p is 1, R² is hydrogen, 6-methyl or 7-methyl, R³ is hydrogen, R⁴ is hydrogen, ethoxycarbonyl or 2-aminoethyl, then R⁵ cannot be hydrogen or methyl;

or a salt, pro-drug or solvate thereof;

According to a further aspect of the third feature of the invention there is provided a compound of Formula (VIc)



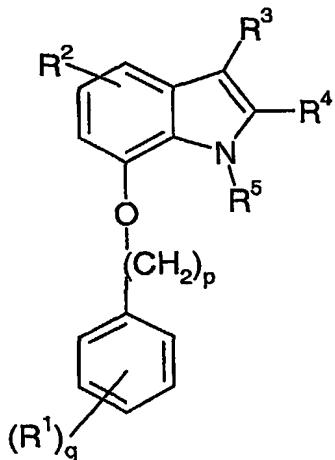
Formula (VIc)

5 wherein: p, q, R¹, R², R³, R⁴, and R⁵ are as defined for Formula (VI) above
with the proviso that

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II);
- (ii) when (R¹)_q is 4-methoxy or 3,4,5-trimethoxy, p is 1, R² is 5-methoxy, R³ is hydrogen, cyanomethyl or 2-aminoethyl, R⁴ is ethoxycarbonyl then R⁵ cannot be 10 hydrogen; and
- (iii) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is methyl, cyanomethyl, 2-aminoethyl, 2-aminopropyl or hydroxyethyl, R⁴ is hydrogen then R⁵ cannot be hydrogen.

or a salt, pro-drug or solvate thereof;

15 According to a further aspect of the third feature of the invention there is provided a compound of Formula (VId)



Formula (VId)

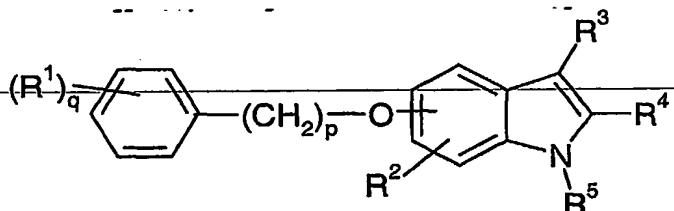
wherein: p, q, R¹, R², R³, R⁴, and R⁵ are as defined for Formula (VI) above
20 with the proviso that

- 12 -

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II); and
- (ii) when q is 0 or (R¹)_q is 4-methoxy or 4-amino, p is 0 or 1 R³ is hydrogen or methyl, R⁴ is hydrogen or ethoxycarbonyl, then R⁵ cannot be hydrogen or methyl.

or a salt, pro-drug or solvate thereof.

5 According to a further aspect of the third feature of the invention there is provided a compound of Formula (VIe)



Formula (VIe)

wherein:

- 10 R¹ is independently selected from hydroxy, amino, alkanoylamino, —OPO₃H₂, or C₁₋₄alkoxy,
wherein the amino group is optionally substituted by an amino acid residue and the
hydroxy group is optionally esterified;
wherein: p, q, R², R³, R⁴, and R⁵ are as defined for Formula (VI) above
with the proviso that

- 15 (i) when R³ is cyano then R⁴ cannot be a group of Formula (II); and
(ii) when (R¹)_q is 4-methoxy, 4-amino or 3,4,5-trimethoxy, p is 0 or 1, R² is hydrogen,
4-chloro or 5-methoxy, R³ is hydrogen, cyanomethyl or 2-aminoethyl, R⁴ is
hydrogen or ethoxycarbonyl, then R⁵ cannot be hydrogen or methyl;
or a salt, pro-drug or solvate thereof.

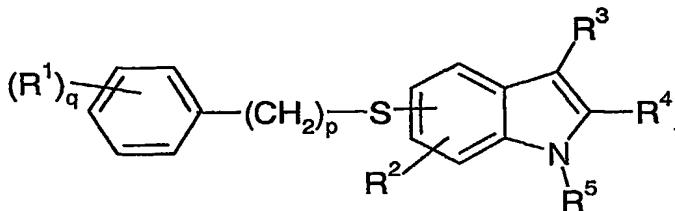
20 According to a further aspect of the third feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (VI), Formula (VIA),
Formula (VIB), Formula (VIC), Formula (VID) or Formula (VIe) or pharmaceutically-
acceptable salt, pro-drug or solvate thereof.

According to a further aspect of the third feature of the invention there is provided a
25 pharmaceutical composition comprising a compound of Formula (VI), Formula (VIA),
Formula (VIB), Formula (VIC), Formula (VID) or Formula (VIe) or a pharmaceutically-
acceptable salt, pro-drug or solvate thereof, in admixture with a pharmaceutically-acceptable
carrier.

According to a further aspect of the third feature of the present invention there is provided the use of a compound of Formula (VI), , Formula (VIa), Formula (VIb), Formula (VIc), Formula (VID) or Formula (VIE), or pharmaceutically-acceptable salt, pro-drug or solvate thereof for the manufacture of a medicament to inhibit and/or reverse and/or alleviate 5 symptoms of angiogenesis and/or any disease state associated with angiogenesis.

According to a further aspect of the third feature of the invention there is provided a method of treatment, in a warm-blooded animal, to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis comprising administering to said warm-blooded animal a therapeutically (including prophylactically) 10 effective amount of a compound of Formula (VI), , Formula (VIa), Formula (VIb), Formula (VIc), Formula (VID) or Formula (VIE), or a pharmaceutically acceptable salt, pro-drug or solvate thereof.

The invention also provides a further novel group of indole compounds. Thus, according to a fourth feature of the invention there is provided a compound of Formula (VII)



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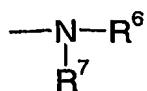
Formula (VII)

wherein:

R^1 is independently selected from halo, hydroxy, amino, alkanoylamino, $-\text{OPO}_3\text{H}_2$, or $\text{C}_{1-4}\text{alkoxy}$, wherein the amino group is optionally substituted by an amino acid residue 20 and the hydroxy group is optionally esterified;

R^2 is selected from: hydrogen, $\text{C}_{1-4}\text{alkyl}$ or $\text{C}_{1-4}\text{alkoxy}$;

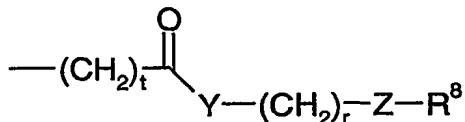
R^3 and R^4 are independently selected from: hydrogen, $\text{C}_{1-4}\text{alkyl}$, $\text{C}_{1-4}\text{alkanoyl}$, $\text{C}_{1-4}\text{alkoxycarbonyl}$, $\text{C}_{1-4}\text{alkoxycarbonylC}_{1-4}\text{alkyl}$, $\text{C}_{1-4}\text{alkoxycarbonylamino}$, amino, amino $\text{C}_{1-4}\text{alkyl}$, carbamoyl, carbamoyl $\text{C}_{1-4}\text{alkyl}$, cyano, cyano $\text{C}_{1-4}\text{alkyl}$, hydroxy, 25 hydroxy $\text{C}_{1-4}\text{alkyl}$; or a group of Formula (II):



Formula (II)

R^6 is hydrogen or C_{1-4} alkyl;

R^5 and R^7 are independently selected from hydrogen, C_{1-4} alkyl or a group of Formula (III):



Formula (III)

5 wherein Y is selected from ---NH--- , ---O--- or a bond;

Z is selected from ---NH--- , ---O--- , ---C(O)--- or a bond;

r is an integer from 0 to 4;

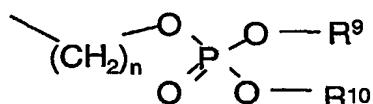
t is an integer from 0 to 1;

10 R^8 is hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy, aryl, 5 or 6 membered heterocyclyl,

5- or 6-membered heteroaryl, wherein aryl, heteroaryl or heterocyclyl

is optionally substituted by C_{1-4} alkyl or C_{1-4} alkoxy, or a group of

Formula (IV):



Formula (IV)

15 wherein n is an integer of from 1 to 6, and;

R^9 and R^{10} are independently selected from hydrogen or
 C_{1-4} alkyl or aryl; and

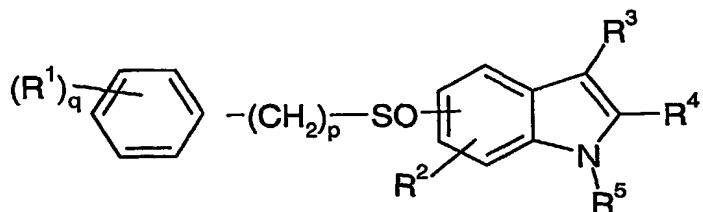
p is an integer between 0 and 1; and

q is an integer between 0 and 3;

20 with the proviso that

- (i) when R^3 is cyano then R^4 cannot be a amino or a group of Formula (II);
- (ii) when q is 0 or $(R^1)_q$ is 4-amino, p is 0 or 1, R^2 is hydrogen, R^3 is hydrogen, R^4 is hydrogen or ethoxycarbonyl, then R^5 cannot be hydrogen;
- (iii) when q is 0, p is 0 or 1, R^2 is hydrogen, R^3 is hydrogen, methyl, cyano, 2-aminoethyl, 1-methyl-2-aminoethyl or 2-aminopropyl, R^4 is hydrogen or amino, then R^5 cannot be hydrogen; and
- (iv) when q is 0, p is 0, R^2 is hydrogen, R^3 is ethoxycarbonyl, then R^5 cannot be hydrogen;

According to a further aspect of the fourth feature of the invention there is provided a compound of Formula (VIIa)



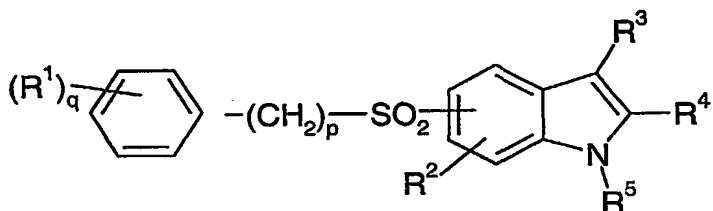
Formula (VIIa)

5 wherein:

wherein: p, q, R¹, R², R³, R⁴, and R⁵ are as defined for Formula (VII) above with the proviso that

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II); or a salt, pro-drug or solvate thereof.

10 According to a further aspect of the fourth feature of the invention there is provided a compound of Formula (VIIb)



Formula (VIIb)

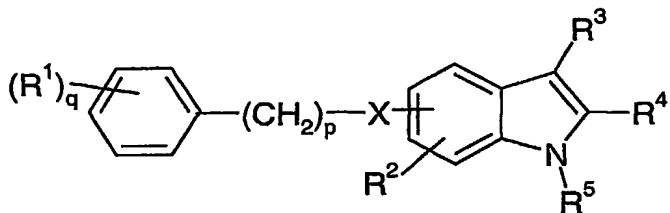
wherein:

15 wherein: p, q, R¹, R², R³, R⁴, and R⁵ are as defined for Formula (VII) above with the proviso that

- (i) when R^3 is cyano then R^4 cannot be a group of Formula (II); and
 - (ii) when $(R^1)_q$ is 4-chloro, p is 0, R^2 is hydrogen, R^3 is hydrogen, R^4 is hydrogen or ethoxycarbonyl, then R^5 cannot be hydrogen;

20 or a salt, pro-drug or solvate thereof.

According to a further aspect of the fourth feature of the invention there is provided a compound of Formula (VIc)



Formula (VIIc)

5 wherein:

X is selected from: —S—, —SO— or —SO₂—;

R^1 is independently selected from hydroxy, amino, alkanoylamino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;

10 q is an integer between 1 and 3;

and wherein: p, R², R³, R⁴, and R⁵ are as defined for Formula (VI) above

with the proviso that

- (i) when R^3 is cyano then R^4 cannot be a group of Formula (II); and
 - (ii) when $(R^1)_q$ is 4-amino, p is 0, R^2 is hydrogen, R^3 is hydrogen, R^4 is hydrogen or ethoxycarbonyl, then R^5 cannot be hydrogen;

or a salt, pro-drug or solvate thereof.

According to a further aspect of the fourth feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (VII), Formula (VIIa), Formula (VIIb) or Formula (VIIc), or pharmaceutically-acceptable salt, or pro-drug solvate

20 thereof.

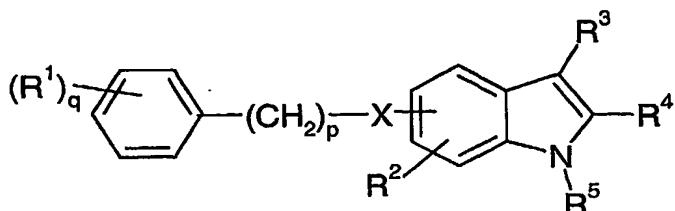
According to a further aspect of the fourth feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (VII), Formula (VIIa) Formula (VIIb) or Formula (VIIc), or a pharmaceutically-acceptable salt, pro-drug or solvate thereof, in admixture with a pharmaceutically-acceptable diluent or carrier.

According to a further aspect of the fourth feature of the present invention there is provided the use of a compound of Formula (VII), Formula (VIIa), Formula (VIIb) or Formula (VIIc), or pharmaceutically acceptable salt, pro-drug or solvate thereof, for the

According to a further aspect of the fourth feature of the invention there is provided a method of treatment, in a warm-blooded animal, to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis comprising administering to said warm-blooded animal a therapeutically (including prophylactically) effective amount of a compound of Formula (VII), Formula (VIIa), Formula (VIIb) or Formula (VIIc), or a pharmaceutically acceptable salt, pro-drug or solvate thereof.

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The invention also provides a further novel group of indole compounds. Thus, according to a fifth feature of the invention there is provided a compound of Formula (VIId)



10

Formula (VIId)

wherein:

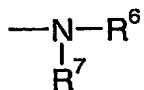
R¹ is independently selected from hydroxy, amino, alkanoylamino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;

15 X is selected from: —O—, —S—, —SO— or —SO₂—;

R² is selected from: hydrogen, C₁₋₄alkyl or C₁₋₄alkoxy;

R³ and R⁴ are independently selected from: hydrogen, C₁₋₄alkyl, C₁₋₄alkanoyl, C₁₋₄alkoxycarbonyl, C₁₋₄alkoxycarbonylC₁₋₄alkyl, C₁₋₄alkoxycarbonylamino, amino, aminoC₁₋₄alkyl, carbamoyl, carbamoylC₁₋₄alkyl, cyano, cyanoC₁₋₄alkyl, hydroxy,

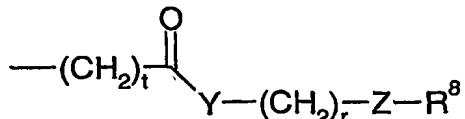
20 hydroxyC₁₋₄alkyl; or a group of Formula (II):



Formula (II)

R⁶ is hydrogen or C₁₋₄alkyl;

R⁵ and R⁷ are independently selected from hydrogen, C₁₋₄alkyl or a group of Formula (III):



Formula (III)

25

wherein Y is selected from —NH—, —O— or a bond;

Z is selected from —NH—, —O—, —C(O)— or a bond;

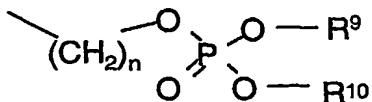
r is an integer from 0 to 4;

t is an integer from 0 to 1;

5 R⁸ is hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy, aryl, 5 or 6 membered heterocyclyl,

5- or 6-membered heteroaryl, wherein aryl, heteroaryl or heterocyclyl
is optionally substituted by C₁₋₄alkyl or C₁₋₄alkoxy, or a group of

Formula (IV):



10 Formula (IV)

wherein n is an integer of from 1 to 6, and;

R⁹ and R¹⁰ are independently selected from hydrogen or
C₁₋₄alkyl or aryl; and

p is an integer between 0 and 1; and

15 q is an integer between 1 and 3;

with the proviso that

(i) when R³ is cyano then R⁴ cannot be a group of Formula (II); and

(ii) when (R¹)_q is 4-methoxy, 4-amino, 4-chloro or 3,4,5-trimethoxy, p is 0 or 1, R² is
hydrogen, 4-chloro or 5-methoxy, R³ is hydrogen, cyanomethyl or 2-aminoethyl,
20 R⁴ is hydrogen or ethoxycarbonyl, then R⁵ cannot be hydrogen or methyl;

or a salt, pro-drug or solvate thereof.

According to a further aspect of the fifth feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (VIIId), or pharmaceutically-acceptable salt, or pro-drug solvate thereof.

25 According to a further aspect of the fifth feature of the invention there is provided a pharmaceutical composition comprising a compound of Formula (VIIId), or a pharmaceutically-acceptable salt, pro-drug or solvate thereof, in admixture with a pharmaceutically-acceptable diluent or carrier.

According to a further aspect of the fifth feature of the present invention there is
30 provided the use of a compound of Formula (VIIId), or pharmaceutically acceptable salt

and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis.

According to a further aspect of the fifth feature of the invention there is provided a method of treatment, in a warm-blooded animal, to inhibit and/or reverse and/or alleviate 5 symptoms of angiogenesis and/or any disease state associated with angiogenesis comprising administering to said warm-blooded animal a therapeutically (including prophylactically) effective amount of a compound of Formula (VIIId), or a pharmaceutically acceptable salt, pro-drug or solvate thereof.

Whilst pharmaceutically acceptable salts of compounds of the invention are preferred, 10 other non-pharmaceutically acceptable salts of compounds of the invention may also be useful, for example in the preparation of pharmaceutically-acceptable salts of compounds of the invention.

For the avoidance of doubt when q is 0, all positions on the phenyl ring are substituted by hydrogen.

15 For the avoidance of doubt the use of the term $(R_1)_q$ when q is between 1 and 3, means that there are 1, 2 or 3 R^1 substituents on the phenyl ring, which when q is 2 or 3 can be the same group or different groups. For example, where $(R_1)_q$ is 3-chloro-4-methoxy then q is 2 and the phenyl ring has a chloro group at the 3-position and a methoxy group at the 4-position, in relation to the $-(CH_2)_pX-$ group, and for example, when $(R_1)_q$ is di-halo, then q is 2 and 20 the phenyl ring has two halo substituents which may be the same group or different groups, wherein the halo groups occupy 2 positions on the phenyl ring.

For the avoidance of doubt the use of the term $(R_1)_{q-1}$ means there is one substituent on the phenyl ring as indicated in the respective formula and the phenyl ring can either bear no further substituents or may bear one or two further substituents. Where there are two further 25 substituents these may be the same group or different groups.

In this specification the generic term 'alkyl' includes both straight-chain and branched-chain alkyl groups. However references to individual alkyl groups such as 'propyl' are specific for the straight-chain version only and references to individual branched-chain alkyl groups such as 'isopropyl' are specific for the branched-chain version only. An 30 analogous convention applies to other generic terms.

The term "aryl" refers to phenyl or naphthyl.

The term "heteroaryl" refers to a 5-10 membered aromatic mono or bicyclic ring containing up to 5 heteroatoms independently selected from nitrogen, oxygen or sulphur,

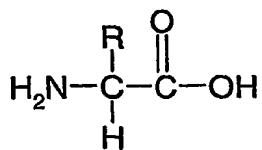
linked via ring carbon atoms or ring nitrogen atoms where a bond from a nitrogen is allowed, for example no bond is possible to the nitrogen of a pyridine ring, but a bond is possible through the 1-nitrogen of a pyrazole ring. Examples of 5- or 6-membered heteroaryl ring systems include pyrrole, furan, imidazole, triazole, pyrazine, pyrimidine, pyridazine, pyridine, 5 isoxazole, oxazole, 1,2,4 oxadiazole, isothiazole, thiazole and thiophene. A 9 or 10 membered bicyclic heteroaryl ring system is an aromatic bicyclic ring system comprising a 6-membered ring fused to either a 5 membered ring or another 6 membered ring. Examples of 5/6 and 6/6 bicyclic ring systems include benzofuran, benzimidazole, benzthiophene, benzthiazole, benzisothiazole, benzoxazole, benzisoxazole, indole, pyridoimidazole, pyrimidoimidazole, 10 quinoline, isoquinoline, quinoxaline, quinazoline, phthalazine, cinnoline and naphthyridine.

The term "heterocyclyl" refers to a 5-10 membered saturated or partially saturated mono or bicyclic ring containing up to 5 heteroatoms selected from nitrogen, oxygen or sulphur linked via ring carbon atoms or ring nitrogen atoms. Examples of 'heterocyclyl' include pyrrolinyl, pyrrolidinyl, morpholinyl, piperidinyl, piperazinyl, dihydropyridinyl and 15 dihydropyrimidinyl

The term **halo** refers to fluoro, chloro, bromo or iodo.

The term "**carbamoyl**" refers to the group $-C(O)-NH_2$.

The term **amino acid residue** is defined as that derived from the coupling of an L-amino acid with an amino group via an amide bond. This bond can either be formed via a 20 carboxylate group on the amino acid backbone or via a side chain carboxylate group, preferably via a carboxylate group on the amino acid backbone. Amino acid residues include those derived from natural and non-natural amino acids, preferably natural amino acids and include α -amino acids β -amino acids and γ -amino acids. For the avoidance of doubt amino acids include those with the generic structure:



25

where R is the amino acid side chain: The definition of amino acid also includes amino acid analogues which have additional methylene groups within the amino acid backbone, for example β -alanine and amino acids which are not naturally occurring such as cyclohexylamine. β -alanine amino acid residue is a naturally occurring amino acid

Preferred amino acids include glycine, alanine, valine, leucine, isoleucine, methionine, proline, phenylalanine, tryptophan, serine, threonine, cysteine, tyrosine, asparagine, glutamine, aspartic acid, glutamic acid, lysine, arginine, histidine, β -alanine and ornithine.

More preferred amino acids include glutamic acid, serine, threonine, arginine, glycine, 5 alanine, β -alanine and lysine. Especially preferred amino acids include glutamic acid, serine, and glycine.

Esterifying groups at R¹ are esterifying groups which increase the solubility of the molecule in water at a pH of approximately pH=7. Such groups include groups with ionisable groups, such as acidic functions or basic functions and groups containing a hydrophilic 10 function. Basic functions include: amino, morpholino, piperidino, piperazino, pyrrolidino, amino acids and imidazolino. Acidic functions include: carboxy, sulphonic acid, phosphate, sulphate and acid mimetics such as tetrazolyl. Hydrophilic groups include hydroxyl.

Suitable R¹ groups wherein hydroxy is esterified include: C₁₋₆alkanoyloxy, arylcarbonyloxy, heterocyclcarbonyloxy, heteroarylcarbonyloxy wherein the R¹ group is 15 optionally substituted with between 1 and 3 groups selected from C₁₋₄alkyl, C₁₋₄alkanoyl, C₁₋₄alkanoylC₁₋₄alkyl, C₁₋₄alkanoylheterocycl, hydroxy, hydroxyC₁₋₄alkyl, carboxy, carboxyphenyl, phosphono, phosphonoC₁₋₄alkyl, amino, aminoC₁₋₄alkyl, N-C₁₋₄alkylamino, N,N-diC₁₋₄alkylamino, carbamoyl, carbamoylC₁₋₄alkyl, heterocycl, heterocyclC₁₋₄alkyl, heterocyclcarbonyl, heterocyclC₁₋₄alkanoylamino, carbamoylheterocycl, [wherein optional 20 substituents comprising heterocycl are optionally further substituted by C₁₋₄alkyl, hydroxyC₁₋₄alkyl, C₁₋₄alkoxyC₁₋₄alkyl, C₁₋₄alkanoyl and formyl, wherein the carbamoyl and amino optional substituents are optionally further N-substituted by C₁₋₄alkyl, di-C₁₋₄alkyl, hydroxyC₁₋₄alkyl, di-(hydroxyC₁₋₄alkyl), carboxyC₁₋₄alkyl, and wherein the amino group is optionally substituted by an amino acid residue] with the proviso that when R₁ is 25 C₁₋₆alkanoyloxy or arylcarbonyloxy R₁ is not unsubstituted and R₁ is not substituted by C₁₋₄alkyl.

More preferred R¹ groups wherein hydroxy is esterified include: carboxypentanoyloxy, 4-carboxyphenylpropanoyloxy; 4-(N-methylpipeazin-1-ylethyl)phenylcarbonyloxy, 4-(piperazin-1-ylethyl)phenylcarbonyloxy, 30 4-[N-di-(hydroxyethyl)aminomethyl]phenylcarbonyloxy, 3-(N-acetylpirazin-1-ylethyl)phenylcarbonyloxy, 3-[N-di-(hydroxyethyl)aminomethyl]phenylcarbonyloxy,

- 4-(N-methylpiperazin-1-ylpropanoylamino)phenylcarbonyloxy,
N-methylpiperazin-1-ylcarbonylpropanoyloxy,
N-di-(hydroxyethyl)aminocarbonylpropanoyloxy, piperazin-1-ylcarbonylpropanoyloxy,
(N-acetyl

5 (N-di-(hydroxyethyl)aminocarbonylpropanoyloxy, and
4-(piperazin-1-ylmethyl)phenylcarbonyloxy.

Further preferred R¹ groups wherein hydroxy is esterified include:

-
- 4-(N-methylpiperazin-1-ylpropanoylamino)phenylcarbonyloxy,
N-methylpiperazin-1-ylcarbonylpropanoyloxy and
- 10 N-di-(hydroxyethyl)aminocarbonylpropanoyloxy.

Examples of C₁₋₄alkyl include methyl, ethyl, propyl, isopropyl, sec-butyl and *tert*-butyl; examples of C₁₋₄alkoxy include methoxy, ethoxy, propoxy and *tert*-butoxy; examples of C₁₋₄alkanoyl include acetyl, propanoyl and butanoyl; examples of C₁₋₄alkanoylamino include acetylamino, propanoylamino and butanoylamino; examples of C₁₋₄alkoxycarbonyl include methoxycarbonyl, ethoxycarbonyl and *tert*-butyloxycarbonyl; examples of C₁₋₄alkoxycarbonylC₁₋₄alkyl include methoxycarbonylethyl, ethoxycarbonylmethyl and *tert*-butyloxycarbonylbutyl; examples of C₁₋₄alkoxycarbonylamino include methoxycarbonylamino, ethoxycarbonylamino and *tert*-butyloxycarbonylamino; examples of aminoC₁₋₄alkyl include aminomethyl, aminoethyl and aminopropyl; examples of 20 cyanoC₁₋₄alkyl include cyanomethyl, cyanoethyl and cyanopropyl; examples of carbamoylC₁₋₄alkyl include carbamoylmethyl, carbamoylethyl and carbamoylpropyl; examples of hydroxyC₁₋₄alkyl include hydroxymethyl, hydroxyethyl and hydroxypropyl.

It is to be understood that, insofar as certain of the compounds in the different features of the invention may exist in optically active or racemic forms by virtue of one or more 25 asymmetric carbon atoms, the invention includes in its definition any such optically active or racemic form which possesses the property of inhibiting and/or reversing and/or alleviating the symptoms of angiogenesis and/or any disease states associated with angiogenesis. The synthesis of optically active forms may be carried out by standard techniques of organic chemistry well known in the art, for example by synthesis from optically active starting 30 materials or by resolution of a racemic form. Similarly, activity of these compounds may be evaluated using the standard laboratory techniques referred to hereinbefore.

The invention also relates to any and all tautomeric forms of the compounds of the different features of the invention that possess the property of inhibiting and/or reversing and/or alleviating the symptoms of angiogenesis and/or any disease states associated with angiogenesis.

- 5 It will also be understood that certain compounds of the present invention may exist in solvated, for example hydrated, as well as unsolvated forms. It is to be understood that the present invention encompasses all such solvated forms which possess the property of inhibiting and/or reversing and/or alleviating the symptoms of angiogenesis and/or any disease states associated with angiogenesis.
- 10 A suitable pharmaceutically-acceptable salt of a compound of the invention is, for example, an acid-addition salt of a compound of the invention which is sufficiently basic, for example, an acid-addition salt with, for example, an inorganic or organic acid, for example hydrochloric, hydrobromic, sulphuric, phosphoric, trifluoroacetic, citric or maleic acid. In addition a suitable pharmaceutically-acceptable salt of a carbazole derivative of the invention 15 which is sufficiently acidic is an alkali metal salt, for example a sodium or potassium salt, an alkaline earth metal salt, for example a calcium or magnesium salt, an ammonium salt or a salt with an organic base which affords a physiologically acceptable cation, for example a salt with methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.
- 20 A preferred group of values of R¹ in each feature of the invention is hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified. A further preferred group of values of R¹ in each feature of the invention is hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue. A more further preferred 25 group of values of R¹ in each feature of the invention is hydroxy, amino, —OPO₃H₂, methoxy, ethoxy, glutamylamino preferably α -glutamylamino, serylamino, glycylamino or alanylarnino. A yet more further preferred group of values of R¹ in each feature of the invention is hydroxy, methoxy, α -glutamylamino, seryl, —OPO₃H₂ or methoxy. A most preferred group of values of R¹ in each feature of the invention is 3,4-dimethoxy, 3,5-dimethoxy, 2,5-dimethoxy, 3,4,5-trimethoxy, 4-hydroxy-3,5-dimethoxy or 30 4-phosphonoxy-3,5-dimethoxy.

A preferred group of values in each feature of the invention for p is 0 or 1. More preferably in each features of the invention p is 0.

A preferred group of values in each feature of the invention for q is 2 or 3. More preferably in each features of the invention q is 3.

5 A preferred group of values of X in the first and second features of the invention is —O—, or —S—. Most preferably X is —S—.

A preferred group of values for R² in each feature of the invention is hydrogen.

A preferred group of values for R³ in each feature of the invention is hydrogen,

C₁₋₄alkanoyl, C₁₋₄alkoxycarbonyl, hydroxyC₁₋₄alkyl, carbamoyl or C₁₋₄alkylcarbamoyl, A
10 more-preferred group of values for R³ in each feature of the invention is cyano,

C₁₋₄alkylcyano, carbamoyl or C₁₋₄alkylcarbamoyl. A most preferred group of values for R³ in
each feature of the invention is C₁₋₄alkylcyano or cyano.

A preferred group of values for R⁴ in each feature of the invention is hydrogen,
carbamoyl, cyano or a group of Formula (II). A more preferred group of values for R⁴ is
15 hydrogen, cyano, carbamoyl, carbamoylC₁₋₄alkyl or C₁₋₄alkoxy. A most preferred value for R⁴
is hydrogen.

A preferred group of values for R⁵ in each aspect of the invention is hydrogen, C₁₋₄alkyl
or a group of Formula (III) wherein t, Y, r, Z and R⁸ are as defined above. A more preferred
group of values for R⁵ in each aspect of the invention is hydrogen or C₁₋₄alkyl. A further
20 preferred value for R⁵ is hydrogen, methyl or ethyl. A most preferred value for R⁵ is methyl.

A preferred group of values for R⁶ in each aspect of the invention is hydrogen or
C₁₋₄alkyl. More preferably R⁶ is hydrogen or methyl. Most preferably R⁶ is hydrogen.

A preferred group of values for R⁷ in each aspect of the invention is hydrogen, methyl,
or a group of Formula (III). A most preferred group of values for R⁷ is hydrogen or a group of
25 Formula (III). A most preferred value of R⁷ is hydrogen.

A preferred group of values for t in each aspect of the invention is 0 or 1. A most
preferred value for t in each aspect of the invention is 0.

A preferred group of values for Y in each aspect of the invention is —O— or a bond.

A most preferred group of values for Y in each aspect of the invention is —O—.

30 A preferred group of values for r in each aspect of the invention is 1, 2 or 3. A most
preferred values for r in each aspect of the invention is 1.

A preferred group of values for Z in each aspect of the invention is —C(O)— or a bond.
A most preferred group of values for Z in each aspect of the invention is a bond.

A preferred group of values for R⁸ in each aspect of the invention is hydrogen, C₁₋₄alkyl C₁₋₄alkoxy, an aryl group, a 6-membered heterocyclic group or a group of Formula (IV),
5 wherein the aryl group and heterocyclic group are optionally substituted by C₁₋₄alkyl or C₁₋₄alkoxy: More preferably R⁸ is a non-aromatic heterocyclic group, an aryl group or a group of Formula (IV), wherein the aryl group and non-aromatic heterocyclic group are optionally substituted by C₁₋₄alkyl or C₁₋₄alkoxy. Further preferably R⁸ is phenyl, morpholino, piperazinyl or a group of Formula (IV), wherein the phenyl, morpholino and piperazinyl group
10 are optionally substituted by methyl or acetyl and R⁹ and R¹⁰ are both hydrogen. Most preferably R⁸ is phenyl, morpholino or 4-methyl-piperazin-1-yl.

A preferred group of values for n in each aspect of the invention is 2 or 3. Preferably n is 2.

A preferred group of values for R⁹ and R¹⁰ in each aspect of the invention is R⁹ and R¹⁰,
15 are independently selected from hydrogen or an alkaryl group, preferably -(CH₂)_m-Ph where m is an integer of from 1 to 6, preferably m is 1. More preferably R⁹ and R¹⁰ are independently selected from hydrogen or benzyl (-CH₂-Ph). Preferably R⁹ and R¹⁰ are both hydrogen.

A preferred group of values for a group of Formula (III) is phenoxy carbonyl,
20 phosphonoxyethylaminocarbonyl, 4-methylpiperazin-1-ylpropoxycarbonyl, carbamoylmethyl, 4-acetyl-piperazin-1-yethoxycarbonyl, 4-methyl-piperazin-1-ylcarbonylpropanoyl and morpholinoethoxycarbonyl. Preferably phenoxy carbonyl, 4-methylpiperazin-1-ylpropoxycarbonyl or carbamoylmethyl.

Preferably only one of R⁵ and R⁷ is a group of Formula (III).

25 In the following preferred groups of compounds of each aspect of the invention values for R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, n, p, q, r, t, X, Y and Z are as hereinbefore defined, unless specifically defined with a preferred group of compounds.

A preferred group of compounds in the first and second features of the invention; comprise compounds wherein:

30 X is —O—;
or salt, pro-drug or solvate thereof.

A preferred group of compounds in the first and second features of the invention; comprise compounds wherein:

X is —S—, —SO— or —S(O)₂—;
or salt, pro-drug or solvate thereof.

5 A further preferred group of compounds in each feature of the invention, comprise compounds wherein:

R³ is selected from cyano or cyanoC₁₋₄alkyl, preferably cyano;
or salt, pro-drug or solvate thereof.

A further preferred group of compounds of each feature of the invention; comprise
10 compounds wherein:

R¹ is amino, hydroxy or —OPO₃H₂, wherein the amino group is optionally substituted by
an amino acid residue and the hydroxy group is optionally esterified;
or salt, pro-drug or solvate thereof.

A further preferred group of compounds of each feature of the invention, comprise
15 compounds wherein:

R¹ is independently selected from hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the
amino group is optionally substituted by an amino acid residue and the hydroxy group
is optionally esterified;

R³ is selected from carbamoyl, carbamoylC₁₋₄alkyl, cyano or cyanoC₁₋₄alkyl; preferably
20 cyano or cyanoC₁₋₄alkyl; and

R⁵ is hydrogen or C₁₋₄alkyl;
or salt, pro-drug or solvate thereof.

A further preferred group of compounds of the first and second features of the
invention, comprise compounds wherein:

25 R¹ is independently selected from hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the
amino group is optionally substituted by an amino acid residue and the hydroxy group
is optionally esterified;

X is —S— or —O—;

R³ is selected from carbamoyl, carbamoylC₁₋₄alkyl, cyano or cyanoC₁₋₄alkyl; preferably
30 cyano or cyanoC₁₋₄alkyl; and

R⁵ is hydrogen or C₁₋₄alkyl;

A further preferred group of compounds in each feature of the invention, comprise compounds wherein:

R¹ is independently selected from hydroxy, —OPO₃H₂, or C₁₋₄alkoxy;

R³ is selected from carbamoyl, carbamoylC₁₋₄alkyl, cyano or cyanoC₁₋₄alkyl; preferably 5 cyano or cyanoC₁₋₄alkyl; and

R⁵ is hydrogen or C₁₋₄alkyl;

or salt, pro-drug or solvate thereof.

A further preferred group of compounds of the first and second features of the invention, comprise compounds wherein:

10 R¹ is independently selected from hydroxy, —OPO₃H₂, or C₁₋₄alkoxy;

X is —S— or —O—;

R³ is selected from carbamoyl, carbamoylC₁₋₄alkyl, cyano or cyanoC₁₋₄alkyl; preferably 15 cyano or cyanoC₁₋₄alkyl; and

R⁵ is hydrogen or C₁₋₄alkyl;

or salt, pro-drug or solvate thereof.

A further preferred group of compounds in each feature of the invention, comprise compounds wherein:

R¹ is selected from hydroxy-dimethoxy, trimethoxy or phosphonoxy-dimethoxy; preferably 20 3,4-dimethoxy, 3,5-dimethoxy, 2,5-dimethoxy, 3,4,5-trimethoxy,

4-hydroxy-3,5-dimethoxy or 4-phosphonoxy-3,5-dimethoxy;

R³ is selected from carbamoyl, carbamoylC₁₋₄alkyl, cyano or cyanoC₁₋₄alkyl; preferably 25 cyano or cyanoC₁₋₄alkyl; and

R⁵ is hydrogen or C₁₋₄alkyl;

or salt, pro-drug or solvate thereof.

25 A further preferred group of compounds of the first and second features of the invention, comprise compounds wherein:

R¹ is selected from hydroxy-dimethoxy, trimethoxy or phosphonoxy-dimethoxy; preferably 30 3,4-dimethoxy, 3,5-dimethoxy, 2,5-dimethoxy, 3,4,5-trimethoxy, 4-hydroxy-3,5-dimethoxy or 4-phosphonoxy-3,5-dimethoxy;

X is —S— or —O—;

R³ is selected from carbamoyl, carbamoylC₁₋₄alkyl, cyano or cyanoC₁₋₄alkyl; preferably 35 cyano or cyanoC₁₋₄alkyl; and

R⁵ is hydrogen or C₁₋₄alkyl;
or salt, pro-drug or solvate thereof.

A further preferred group of compounds of the first and second features of the invention comprise compounds, wherein:

- 5 R¹ is selected from hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue;
- R² is hydrogen;
- X is selected from: —O—, or —S—;
- p is 0 or 1;
- 10 q is an integer between 1 and 3;
- R³ is selected from: hydrogen, cyano, carbamoyl, carbamoylC₁₋₄alkyl, C₁₋₄alkanoy, C₁₋₄alkoxycarbonyl;
- R⁴ is selected from: hydrogen, cyano or carbamoyl;
- R⁵ is hydrogen or C₁₋₄alkyl
- 15 or salt, pro-drug or solvate thereof.

Particular compounds of each feature of the invention include:

- 3-cyano-5-phenylsulphonyl-1*H*-indole;
- 3-cyano-5-phenoxy-1*H*-indole;
- 3-cyano-5-(4-hydroxyphenoxy)-1*H*-indole;
- 20 2-cyano-5-benzyloxy-1*H*-indole;
- 3-Carbamoyl-5-phenoxy-1*H*-indole; and
- 3-Carbamoyl-5-(4-hydroxyphenoxy)-1*H*-indole.

or salt, pro-drug or solvate thereof

More particular compounds of each feature of the invention include:

- 25 1-methyl-3-cyano-5-(4-hydroxy-3,5-dimethoxyphenoxy)-1*H*-indole;
- 1-methyl-3-cyano-5-(4-phosphonoxy-3,5-dimethoxyphenoxy)-1*H*-indole;
- 3-cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole;
- 1-methyl-3-cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole; and
- 1-Methyl-3-cyano-5-(4-hydroxy-3,5-dimethoxyphenoxy)-1*H*-indole.

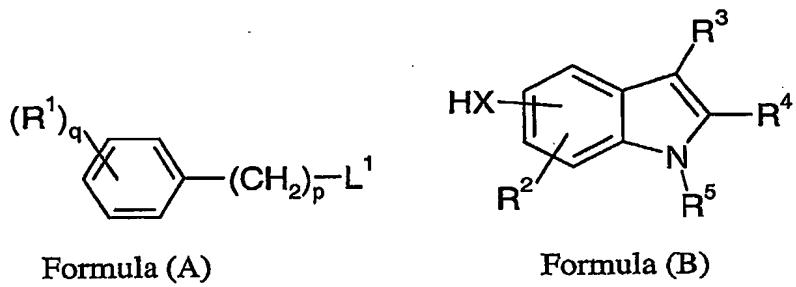
30 or salt, pro-drug or solvate thereof

A compound of the invention or a pharmaceutically-acceptable salt, or solvate thereof,

related compounds. Such processes, when used to prepare a compound of the invention or a pharmaceutically-acceptable salt, or solvate thereof, are provided as a further feature of the invention and are illustrated by the following representative examples in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, X, Y, Z, n, p, q, r and t have the same meaning as herein before defined. The reader is referred to Advanced Organic Chemistry, 4th Edition, by Jerry March, published by John Wiley & Sons 1992, for general guidance on reaction conditions and reagents. The reader is referred to Protective Groups in Organic Synthesis 2nd Edition, by Green et al, published by John Wiley & Sons for general guidance on protecting groups.

Thus, according to the sixth aspect of the invention there is provided a process for preparing a compound of Formula (I), or salt, solvate or pro-drug thereof, which process (wherein R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹, X, Y, Z, n, p, q, r and t are unless otherwise specified as defined in Formula (I)) comprising:

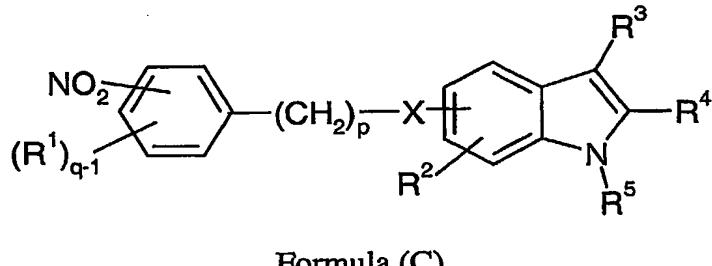
a) for compounds of Formula (I) wherein X is —O—, or —S—, reacting a compound of Formula (A) with a compound of Formula (B),



15

wherein L¹ is a leaving group;

b) for compounds of Formula (I) in which R¹ is amino; reduction of a compound of Formula (C):

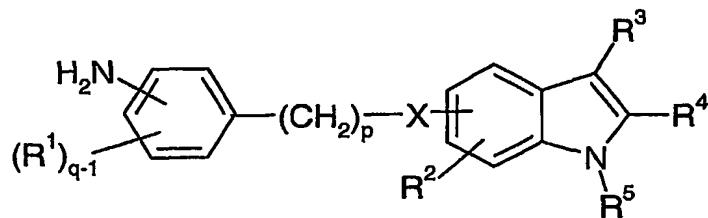


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c) for compounds of Formula (I) wherein R⁵ is C₁₋₄alkyl, reacting a compound of Formula (I) wherein R⁵ is hydrogen with a suitable alkylhalide,

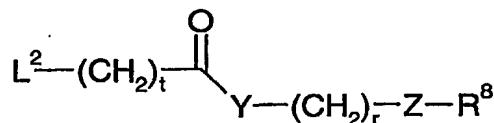
d) for compounds of Formula (I) wherein R¹ is an amino group substituted by an amino acid residue, reacting a compound of Formula (D) with an amino acid,

25



Formula (D)

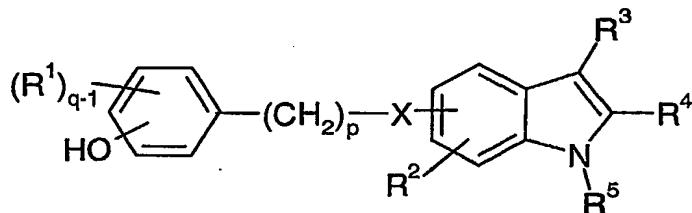
- e) for compounds of Formula (I) in which R^5 is a group of Formula (II) and R^7 is a group of Formula (III), reacting a compounds of Formula (I) in which R^5 is a group of Formula (II) and R^7 is hydrogen with compounds of Formula (E) below, in which L^2 is a leaving group:



Formula (E)

- 10 f) for compounds of Formula (I) in which R^4 is hydrogen, reacting a compounds of Formula (I) in which R^3 is hydrogen and R^4 is hydrogen with a compounds of L^3R^3 in which L^3 is a leaving group;

g) for compounds of Formula (I) in which R^1 is an esterified hydroxyl group, reacting a compound of Formula (F) with an appropriate carboxylic acid or carboxylic acid derivative;



Formula (F)

and thereafter if necessary:

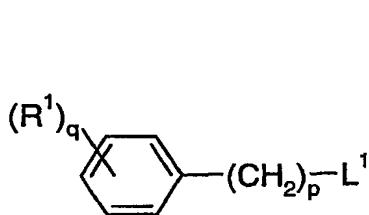
- i) converting a compound of Formula (I) into another compound of Formula (I);
ii) removing any protecting groups;
20 iii) forming a salt, pro-drug or solvate.

According to a further feature of the sixth aspect of the invention there is provided the processes a), b), c), d), e), f) and g) described above for the preparation of compounds of the formula I, or a salts, one, two or mixture thereof.

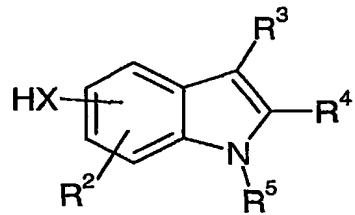
- Process a)* Compounds of Formula (A) and compound of Formula (B) can be reacted together in an organic solvent, at a temperature between room temperature and about 80°C, optionally in the presence of a base such as sodium hydride, potassium carbonate or triethylamine.
- 5 *Process b)* The conditions for reduction of a compound of Formula (C) are well known in the art. Examples of reducing agents include hydrogen and a hydrogenation catalyst (for example palladium on carbon), iron and acetic acid, and zinc and hydrochloric acid. The reaction is preferably carried out in the presence of a suitable solvent such as an alcohol, for example methanol or ethanol, and at a temperature in the range of 0-80°C, preferably at or
10 near room temperature. Further examples of reducing conditions include sodium dithionite in the presence of a base, preferably sodium bicarbonate in a suitable solvent such as DMF or N-methyl-pyrrolidone.

Compounds of Formula (C) can be prepared by reaction of a compound of Formula (F) and a compound of Formula (G) as described in process a).

15



Formula (F)



Formula (G)

- Process c)* Compounds of Formula (I) wherein R₂ is hydrogen and a suitable alkyl halide may be reacted together in a suitable organic solvent such as DMF or DMSO, in the presence of a base, such as sodium hydride or potassium carbonate at a temperature between about
20 room temperature and about 80°C.

- Process d)* Compound of Formula (D) can be reacted with an amino acid using a suitable amide bond forming reaction. Amide bond forming reactions are well known in the art, for example, a carbodiimide coupling reaction can be performed with EDAC in the presence of DMAP in a suitable solvent such as methylene chloride, chloroform or DMF at room
25 temperature.

- Process e)* Compounds of Formula (I) in which R³ is hydrogen can be reacted with compounds of Formula (E) under conditions well known in the art. For example L² may be chloro or p-nitrophenoxy. When L² is Chloro this is carried out in the presence of a base, preferably pyridine.

Process f) Compounds of Formula (I) in which R³ is hydrogen and R⁴ is hydrogen can be reacted with compounds of the formula L³R³ in acetonitrile or diethyl ether in the presence of a base such as triethylamine at a temperature between 0°C and room temperature;

Process g) Processes for the formulation of an ester between a hydroxyl group and a carboxylic acid or carboxylic acid derivative are well known in the art. For example this reaction an acid chloride can be reacted with an alcohol in the presence of a base such as triethyleamine. A carboxylic acid derivative is any derivative of a carboxylic acid which when reacted with a hydroxyl under appropriate conditions will form an ester bond.

Examples of carboxylic acid derivatives include an acid chloride.

10 The compounds used as starting points for the reactions described above are commercially available or they are known compounds or they are prepared by processes known in the art.

It will also be appreciated that in some of the reactions mentioned herein it may be necessary/desirable to protect any sensitive groups in the compounds. The instances where 15 protection is necessary or desirable and suitable methods for protection are known to those skilled in the art. Conventional protecting groups may be used in accordance with standard practice (for illustration see T.W. Green, Protective Groups in Organic Synthesis, John Wiley and Sons, 1991). Thus, if reactants include groups such as amino, carboxy or hydroxy it may be desirable to protect the group in some of the reactions mentioned herein.

20 A suitable protecting group for an amino or alkylamino group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an alkoxycarbonyl group, for example a methoxycarbonyl, ethoxycarbonyl or *tert*-butoxycarbonyl group, an arylmethoxycarbonyl group, for example benzyloxycarbonyl, or an aroyl group, for example benzoyl. The deprotection conditions for the above protecting groups necessarily vary with the choice of 25 protecting group. Thus, for example, an acyl group such as an alkanoyl or alkoxycarbonyl group or an aroyl group may be removed for example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an acyl group such as a *tert*-butoxycarbonyl group may be removed, for example, by treatment with a suitable acid as hydrochloric, sulphuric or phosphoric acid or trifluoroacetic acid and an 30 arylmethoxycarbonyl group such as a benzyloxycarbonyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon, or by treatment with a Lewis

primary amino group is, for example, a phthaloyl group which may be removed by treatment with an alkylamine, for example dimethylaminopropylamine, or with hydrazine.

A suitable protecting group for a hydroxy group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an aroyl group, for example benzoyl, or an arylmethyl group, for example benzyl. The deprotection conditions for the above protecting groups will necessarily vary with the choice of protecting group. Thus, for example, an acyl group such as an alkanoyl or an aroyl group may be removed, for example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an arylmethyl group such as a benzyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

A suitable protecting group for a carboxy group is, for example, an esterifying group, for example a methyl or an ethyl group which may be removed, for example, by hydrolysis with a base such as sodium hydroxide, or for example a *tert*-butyl group which may be removed, for example, by treatment with an acid, for example an organic acid such as trifluoroacetic acid, or for example a benzyl group which may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

The protecting groups may be removed at any convenient stage in the synthesis using conventional techniques well known in the chemical art.

In order to use a compound of the Formula (I), or a pharmaceutically-acceptable salt or 20 in vivo cleavable ester thereof, for the therapeutic treatment (including prophylactic treatment) of mammals including humans, it is normally formulated in accordance with standard pharmaceutical practice as a pharmaceutical composition.

The compositions of the invention may be in a form suitable for oral use (for example as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible 25 powders or granules, syrups or elixirs), for topical use (for example as creams, ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for example as a finely divided powder or a liquid aerosol), for administration by insufflation (for example as a finely divided powder) or for parenteral administration (for example as a sterile aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing 30 or as a suppository for rectal dosing).

The compositions of the invention may be obtained by conventional procedures using conventional pharmaceutical excipients, well known in the art. Thus, compositions intended for oral use may contain, for example, one or more colouring, sweetening, flavouring and/or

preservative agents.

Suitable pharmaceutically-acceptable excipients for a tablet formulation include, for example, inert diluents such as lactose, sodium carbonate, calcium phosphate or calcium carbonate, granulating and disintegrating agents such as corn starch or algenic acid; binding agents such as starch; lubricating agents such as magnesium stearate, stearic acid or talc; preservative agents such as ethyl or propyl p-hydroxybenzoate, and anti-oxidants, such as ascorbic acid. Tablet formulations may be uncoated or coated either to modify their disintegration and the subsequent absorption of the active ingredient within the gastrointestinal tract, or to improve their stability and/or appearance, in either case, using conventional coating agents and procedures well known in the art.

Compositions for oral use may be in the form of hard gelatin capsules in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules in which the active ingredient is mixed with water or an oil such as peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions generally contain the active ingredient in finely powdered form together with one or more suspending agents, such as sodium carboxymethylcellulose, methylcellulose, hydroxypropylmethylcellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents such as lecithin or condensation products of an alkylene oxide with fatty acids (for example polyoxyethylene stearate), or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives (such as ethyl or propyl p-hydroxybenzoate, anti-oxidants (such as ascorbic acid), colouring agents, flavouring agents, and/or sweetening agents (such as sucrose, saccharine or aspartame).

Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil (such as arachis oil, olive oil, sesame oil or coconut oil) or in a mineral oil (such as liquid paraffin). The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set out above, and flavouring agents may be added to provide a palatable oral preparation. These compositions may be prepared

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water generally contain the active ingredient together with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional 5 excipients such as sweetening, flavouring and colouring agents, may also be present.

The pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, such as olive oil or arachis oil, or a mineral oil, such as for example liquid paraffin or a mixture of any of these. Suitable emulsifying agents may be, for example, naturally-occurring gums such as gum acacia or gum 10 tragacanth, naturally-occurring phosphatides such as soya bean, lecithin, an esters or partial esters derived from fatty acids and hexitol anhydrides (for example sorbitan monooleate) and condensation products of the said partial esters with ethylene oxide such as polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening, flavouring and preservative agents.

15 Syrups and elixirs may be formulated with sweetening agents such as glycerol, propylene glycol, sorbitol, aspartame or sucrose, and may also contain a demulcent, preservative, flavouring and/or colouring agent.

The pharmaceutical compositions may also be in the form of a sterile injectable aqueous or oily suspension, which may be formulated according to known procedures using 20 one or more of the appropriate dispersing or wetting agents and suspending agents, which have been mentioned above. A sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example a solution in 1,3-butanediol.

Suppository formulations may be prepared by mixing the active ingredient with a 25 suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Suitable excipients include, for example, cocoa butter and polyethylene glycols.

Topical formulations, such as creams, ointments, gels and aqueous or oily solutions or suspensions, may generally be obtained by formulating an active ingredient with a 30 conventional, topically acceptable, vehicle or diluent using conventional procedures well known in the art.

Compositions for administration by insufflation may be in the form of a finely divided powder containing particles of average diameter of, for example, 30 μm or much less, the

powder itself comprising either active ingredient alone or diluted with one or more physiologically acceptable carriers such as lactose. The powder for insufflation is then conveniently retained in a capsule containing, for example, 1 to 50mg of active ingredient for use with a turbo-inhaler device, such as is used for insufflation of the known agent sodium

5 cromoglycate.

Compositions for administration by inhalation may be in the form of a conventional pressurised aerosol arranged to dispense the active ingredient either as an aerosol containing finely-divided solid or liquid droplets. Conventional aerosol propellants such as volatile fluorinated hydrocarbons or hydrocarbons may be used and the aerosol device is conveniently

10 arranged to dispense a metered quantity of active ingredient.

For further information on formulation the reader is referred to Chapter 25.2 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The amount of active ingredient that is combined with one or more excipients to

15 produce a single dosage form will necessarily vary depending upon the host treated and the particular route of administration. For example, a formulation intended for oral administration to humans will generally contain, for example, from 0.5 mg to 2 g of active agent compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition. Dosage unit forms will

20 generally contain about 1 mg to about 500 mg of an active ingredient. For further information on Routes of Administration and Dosage Regimes the reader is referred to Chapter 25.3 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The size of the dose for therapeutic or prophylactic purposes of a compound of the

25 Formula (I) will naturally vary according to the nature and severity of the conditions, the age and sex of the animal or patient and the route of administration, according to well known principles of medicine.

In using a compound of the Formula (I) for therapeutic or prophylactic purposes it will generally be administered so that a daily dose in the range of, for example, 0.5 mg to 75 mg

30 per kg body weight is received, given if required in divided doses. In general lower doses will be administered when a parenteral route is employed. Thus, for example, for intravenous

ADMINISTRATION OF DRUGS IN THE PRACTICE OF PHARMACEUTICAL SCIENCE, 1E AND 2E EDITIONS BY R. H. WESTON

generally be used. Intranvenous administration is however preferred, typically, intravenous doses of about 10 mg to 500 mg per patient of a compound of this invention.

The compounds of this invention may be used in combination with other drugs and therapies used to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis. Examples of such disease states include: cancer, diabetes, psoriasis, rheumatoid arthritis, Kaposi's sarcoma, haemangioma, acute and chronic nephropathies, atheroma, arterial restenosis, autoimmune diseases, acute inflammation, endometriosis, dysfunctional uterine bleeding and ocular diseases with retinal vessel proliferation.

If formulated as a fixed dose such combination products employ the compounds of this invention within the dosage range described herein and the other pharmaceutically-active agent within its approved dosage range. Sequential use is contemplated when a combination formulation is inappropriate.

In the field of medical oncology examples of such combinations include combinations with the following categories of therapeutic agent:

i) anti-angiogenic agents that work by different mechanisms from the compounds of Formula (I) (for example linomide, inhibitors of integrin $\alpha v\beta 3$ function, angiostatin, endostatin, razoxane, thalidomide) and including vascular endothelial growth factor (VEGF) receptor tyrosine kinase inhibitors (RTKIs) (for example those described in international patent applications publication nos. WO-97/22596, WO-97/30035, WO-97/32856 and WO-98/13354, the entire disclosure of which documents is incorporated herein by reference);

ii) cytostatic agents such as anti-oestrogens (for example tamoxifen, toremifene, raloxifene, droloxifene, iodoxyfene), progestogens (for example megestrol acetate), aromatase inhibitors (for example anastrozole, letrozole, vorazole, exemestane), anti-progestogens, anti-androgens (for example flutamide, nilutamide, bicalutamide, cyproterone acetate), LHRH agonists and antagonists (for example goserelin acetate, luprolide), inhibitors of testosterone 5 α -dihydroreductase (for example finasteride), anti-invasion agents (for example metalloproteinase inhibitors like marimastat and inhibitors of urokinase plasminogen activator receptor function) and inhibitors of growth factor function, (such growth factors include for example epidermal growth factor (EGF), platelet derived growth factor and

- hepatocyte growth factor such inhibitors include growth factor antibodies, growth factor receptor antibodies, tyrosine kinase inhibitors and serine/threonine kinase inhibitors);
- iii) biological response modifiers (for example interferon);
 - iv) antibodies (for example edrecolomab); and
- 5 v) anti-proliferative/anti-neoplastic drugs and combinations thereof, as used in medical oncology, such as anti-metabolites (for example anti-folates like methotrexate, fluoropyrimidines like 5-fluorouracil, purine and adenosine analogues, cytosine arabinoside);
-
- anti-tumour antibiotics (for example anthracyclines like doxorubicin, daunomycin, epirubicin and idarubicin, mitomycin-C, dactinomycin, mithramycin); platinum derivatives (for example
- 10 cisplatin, carboplatin); alkylating agents (for example nitrogen mustard, melphalan, chlorambucil, busulphan, cyclophosphamide, ifosfamide, nitrosoureas, thiotepa); anti-mitotic agents (for example vinca alkaloids like vincristine and taxoids like taxol, taxotere); enzymes (for example asparaginase); thymidylate synthase inhibitors (for example raltitrexed); topoisomerase inhibitors (for example epipodophyllotoxins like etoposide and teniposide,
- 15 amsacrine, topotecan, irinotecan).

The compounds of the invention may also be used in combination with surgery or radiotherapy.

- According to the seventh feature of the present invention there is provided a compound of Formula (I), or salt, solvate or thereof, preferably in the form of a
- 20 pharmaceutical composition, which when dosed in divided doses (also known as split doses) produces a greater anti-tumour effect than when a single dose is given.

Anti-tumour effects include but are not limited to, inhibition of tumour growth, tumour growth delay, regression of tumour, shrinkage of tumour, increased time to re-growth of tumour on cessation of treatment, slowing of disease progression. It is expected that when

25 a compound of the present invention is administered to a warm-blooded animal such as a human, in need of treatment for cancer involving a solid tumour, said method of treatment will produce an effect, as measured by, for example, one or more of: the extent of the anti-tumour effect, the response rate, the time to disease progression and the survival rate.

- According to a further aspect of the seventh feature of the present invention there is
- 30 provided a method for the production of a vascular damaging effect in a warm-blooded animal such as a human which comprises administering to said animal in divided doses an effective

amount of a compound of Formula (I), or salt, or solvate thereof, preferably in the form of a pharmaceutical composition.

According to a further aspect of the seventh feature of the present invention there is provided a method for the treatment of a cancer involving a solid tumour in a warm-blooded animal such as a human, which comprises administering to said animal in divided doses an effective amount of a compound of Formula (I), or salt, or solvate thereof, preferably in the form of a pharmaceutical composition.

According to a further aspect of the seventh feature of the present invention there is provided a medicament comprising two or more fractions of doses of a compound of Formula (I), or salt, or solvate thereof, preferably in the form of a pharmaceutical composition, which together add up to a total daily dose, for administration in divided doses for use in a method of treatment of a human or animal body by therapy.

According to a further aspect of the seventh feature of the present invention there is provided a kit comprising two or more fractions of doses of a compound of Formula (I), or salt, or solvate thereof, preferably in the form of a pharmaceutical composition, which together add up to a total daily dose, for administration in divided doses.

According to a further aspect of the seventh feature of the present invention there is provided a kit comprising:

- a) two or more fractions of doses of a compound of Formula (I), or salt, or solvate thereof, which together add up to a total daily dose, in unit dosage forms for administration in divided doses; and
- b) container means for containing said dosage forms.

According to a further aspect of the seventh feature of the present invention there is provided a kit comprising:

- a) two or more fractions of doses of a compound of Formula (I), or salt, or solvate thereof, which together add up to a total daily dose, together with an excipient or carrier, in unit dosage forms; and
- b) container means for containing said dosage forms.

According to a further aspect of the seventh feature of the present invention there is provided the use of a compound of Formula (I), or salt, or solvate thereof, in the manufacture of a medicament for administration in divided doses for use in the production of a vascular damaging effect in a warm-blooded animal such as a human.

According to a further aspect of the seventh feature of the present invention there is provided the use of a compound of Formula (I), or salt, or solvate thereof, in the manufacture of a medicament for administration in divided doses for use in the production of an anti-cancer effect in a warm-blooded animal such as a human.

5 According to a further aspect of the seventh feature of the present invention there is provided the use of a compound of Formula (I), or salt, or solvate thereof, in the manufacture of a medicament for administration in divided doses for use in the production of an anti-tumour effect in a warm-blooded animal such as a human.

Divided doses, also called split doses, means that the total dose to be administered
10 to a warm-blooded animal, such as a human, in any one day period (for example one 24 hour period from midnight to midnight) is divided up into two or more fractions of the total dose and these fractions are administered with a time period between each fraction of about greater than 0 hours to about 10 hours, preferably about 1 hour to about 6 hours, more preferably about 2 hours to about 4 hours. The fractions of total dose may be about equal or unequal.

15 Preferably the total dose is divided into two parts which may be about equal or unequal.

The time intervals between doses may be for example selected from:
about 1 hour, about 1.5 hours, about 2 hours, about 2.5 hours, about 3 hours, about 3.5 hours,
about 4 hours, about 4.5 hours, about 5 hours, about 5.5 hours and about 6 hours.

20 The time intervals between doses may be any number (including non-integers) of minutes between greater than 0 minutes and 600 minutes, preferably between 45 and 375 minutes inclusive. If more than two doses are administered the time intervals between each dose may be about equal or unequal.

25 Preferably two doses are given with a time interval in between them of greater than or equal to 1 hour and less than 6 hours.

More preferably two doses are given with a time interval in between them of greater than or equal to two hours and less than 5 hours.

Yet more preferably two doses are given with a time interval in between them of greater than or equal to two hours and less than or equal to 4 hours.

30 Particularly the total dose is divided into two parts which may be about equal or unequal with a time interval between doses of greater than or equal to about two hours and

More particularly the total dose is divided into two parts which may be about equal with a time interval between doses of greater than or equal to about two hours and less than or equal to about 4 hours.

For the avoidance of doubt the term 'about' in the description of time periods means 5 the time given plus or minus 15 minutes, thus for example about 1 hour means 45 to 75 minutes, about 1.5 hours means 75 to 105 minutes. Elsewhere the term 'about' has its usual dictionary meaning.

Although the compounds of the Formula (I) are primarily of value as therapeutic agents for use in warm-blooded animals (including man), they are also useful whenever it is 10 required to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis. Thus, they are useful as pharmacological tools for use in the development of new biological tests and in the search for new pharmacological agents.

15 Biological Assay

Colchicine Binding site competitive assay kit.

The ability of a ligand to bind specifically to the colchicine binding site on tubulin, an indicator of the vascular damaging activity, was assessed using a size exclusion chromatography assay kit from "Cytoskeleton" (1650 Fillmore St. #240, Denver, CO 80206, 20 U.S.A.) Catalogue number of kit: BK023.

The following reagents were used:

tubulin buffer, to give 0.1mM GTP, 0.5mM MgCl₂, 0.5mM EGTA, 40mM PIPES buffer at pH 6.9 in the final reaction mix;

purified tubulin protein from bovine brain at 1mg/ml in tubulin buffer;

25 0.02mM fluorescent colchicine in tubulin buffer [FITC (fluorescein isothiocyanate)-labelled];

2mM colchicine in tubulin buffer;

0.2mM vinblastine in tubulin buffer; and

G-25 Sephadex™ Fine - particle size 34-138μm.

30 The reaction was performed as follows:

8 μ l of test compound (dissolved in DMSO) was gently mixed with 150 μ l of tubulin. This was then incubated at 37°C for 30 minutes. Then 4 μ l of the fluorescent colchicine was added, the incubation mix vortexed for 5 seconds and then incubated for a further 30 minutes at 37°C.

At the end of the reaction incubation size exclusion chromatography was performed to

5 separate the tubulin with fluorescent colchicine bound from the free, unbound colchicine. If a test compound inhibited fluorescent colchicine binding then a reduced signal is measured and the compound is confirmed as a colchicine site binding moiety.

Chromatography was performed as follows, using chromatography columns filled with 3mls of G-25 Sephadex™ Fine slurry. The incubation mixture was pipetted onto the column

10 and up to 12 elutions of 160 μ l were collected. The fluorescence of the tubulin-containing fractions was detected on a spectrophotometer which excites at 485nm and emits at 535nm. Control incubations were also performed, 8 μ l DMSO (negative control) and 8 μ l colchicine stock (positive competition control), instead of the 8 μ l of test compound in the incubation mixture.

15 The degree of competition of colchicine binding by either unlabelled colchicine or test compound was calculated relative to the DMSO negative control.

Compounds of Formula (I) encompass vascular damaging agents and pro-drugs of vascular damaging agents. Pro-drugs of vascular damaging agents are believed to be cleaved *in-vivo*. Without being bound by theoretical considerations these pro-drugs may have lower

20 activity in the *in-vitro* colchicine binding site competitive assay, than would be anticipated when the activity of these compounds is measured in cell based assays or *in-vivo*.

The invention will now be illustrated with the following non-limiting Examples in which, unless otherwise stated:

- (i) evaporation were carried out by rotary evaporation in vacuo and work-up;
- 25 procedures were carried out after removal of residual solids such as drying agents by filtration;
- (ii) operations were carried out at ambient temperature, that is in the range 18-25°C and under an atmosphere of an inert gas such as argon or nitrogen;
- (iii) yields are given for illustration only and are not necessarily the maximum attainable;
- 30 (iv) the structures of the end-products of the Formula I were confirmed by nuclear (generally proton) magnetic resonance (NMR) and mass spectral techniques; proton magnetic

shown as follows: s, singlet; d, doublet; t, triplet; m, multiplet; br, broad; q, quartet, quin, quintet;

(v) intermediates were not generally fully characterised and purity was assessed by thin layer chromatography (TLC), high-performance liquid chromatography (HPLC), infra-red 5 (IR) or NMR analysis;

(vi) flash chromatography was performed on silica (Merck Keiselgel: Art.9385);

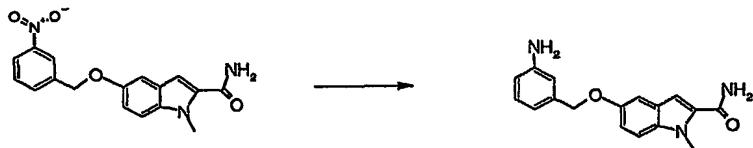
(vii) OASIS™ is a macroporous co-polymer, used to purify hydrophilic compounds, made from a balanced ratio of lipophilic divinylbenzene and hydrophilic N-vinylpyrrolidone. OASIS™ is described in the following patents, US Patent Number 10 No.5882521, US Patent Number No.5976376 and US Patent Number No.6106721. OASIS™ sample extraction products were obtained from Waters Corporation (Milford, Massachusetts, USA).

Abbreviations

15	4-Dimethylaminopyridine	DMAP
	1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride	EDCI
	Di-isopropyl ethylamine	DIEA
	Dimethyl sulphoxide	DMSO
	N-(9-fluorenylmethoxycarbonyl)	N-FMOC
20	N-methylpyrrolidine	NMP

Example 1

1-Methyl-2-carbamoyl-5-(3-aminobenzyloxy)-1*H*-indole



Example 1

25 A suspension of 1-methyl-2-carbamoyl-5-(3-nitrobenzyloxy)-1*H*-indole (1.22 g ; 3.75 mmol), Pt O₂ (0.18 g) and K₂CO₃ (0.18 g) in methanol (120 ml) was hydrogenated under 45 psi for 1 hour.

The mixture was diluted with methanol, filtered on celite. After evaporation the residue was purified by flash chromatography eluting with CH₂Cl₂/CH₃CN/MeOH 46/46/8 to give 1-

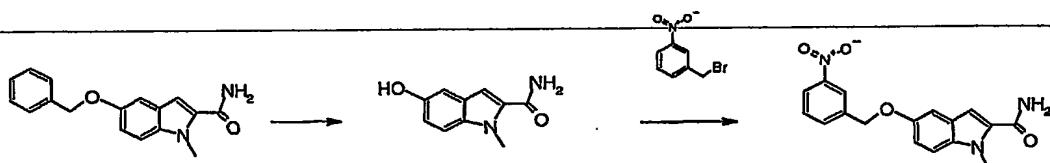
30 methyl-2-carbamoyl-5-(3-aminobenzyloxy)-1*H*-indole.

Yield : 70 %

¹H NMR spectrum (DMSO_d₆) : 3.95 (s, 3H) ; 4.95 (s, 2H) ; 5.08 (bs, 2H) ; 6.49 (ddd, 1H) ; 6.58 (d, 1H) ; 6.65 (dd, 1H) ; 6.94-7.04 (m, 3H) ; 7.13 (d, 1H) ; 7.29 (bs, 1H) ; 7.42 (d, 1H) ; 7.88 (bs, 1H).

5 MS - ESI : 296 [M+H]⁺

The starting material was prepared as follows:



1-Methyl-2-carbamoyl-5-hydroxy-1H-indole

- 10 A suspension of 1-methyl-2-carbamoyl-5-benzyloxy-1*H*-indole (1.94 g ; 6.96 mmol) and 10 % Pd/C (0.5 g) in methanol (100 ml) was hydrogenated for 2 hours. After filtration of the catalyst on celite, the filtrate was evaporated to give 1-methyl-2-carbamoyl-5-hydroxy-1*H*-indole.

Yield : 95 %

- 15 ¹H NMR spectrum (DMSO_d₆) : 3.92 (s, 3H) ; 6.79 (dd, 1H) ; 6.88 (d, 1H) ; 6.92 (s, 1H) ; 7.25 (bs, 1H) ; 7.31 (d, 1H) ; 7.83 (bs, 1H) ; 8.88 (bs, 1H).

1-Methyl-2-carbamoyl-5-(3-nitrobenzyloxy)-1H-indole

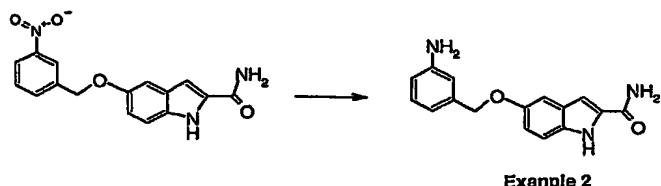
- A mixture of 1-methyl-2-carbamoyl-5-hydroxy-1*H*-indole (0.875 g ; 4.6 mmol), 2-nitrobenzyl bromide (1 g ; 4.63 mmol) and Cs₂CO₃ (1.5 g ; 4.63 mmol) in acetonitrile (55 ml) was heated at 90°C under argon atmosphere for 2 h 30. After evaporation to dryness, the residue was extracted with CH₂Cl₂ / H₂O. The organic phase was evaporated and the residue triturated in CH₂Cl₂ / ether to give 1-methyl-2-carbamoyl-5-(3-nitrobenzyloxy)-1*H*-indole as a solid.

Yield : 86 %

- 25 ¹H NMR spectrum (DMSO_d₆) : 3.96 (s, 3H) ; 5.30 (s, 2H) ; 7.03 (s, 1H) ; 7.05 (dd, 1H) ; 7.21 (d, 1H) ; 7.31 (bs, 1H) ; 7.46 (d, 1H) ; 7.71 (dd, 1H) ; 7.90 (bs, 1H) ; 7.94 (d, 1H) ; 8.19 (dd, 1H) ; 8.33 (s, 1H).

MS - ESI : 326 [M+H]⁺

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2-Carbamoyl-5-(3-aminobenzylxy)-1*H*-indole

2-Carbamoyl-5-(3-aminobenzylxy)-1*H*-indole was prepared using the same general method as described for Example 1.

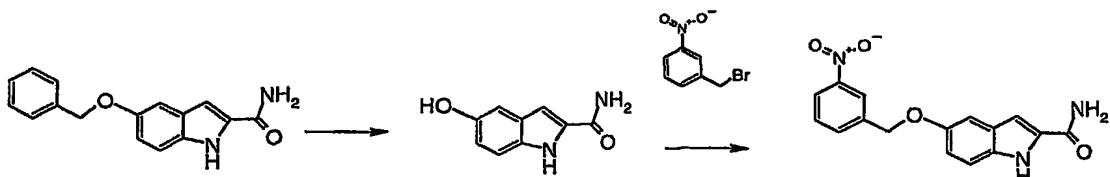
5 Yield : 85 %

¹H NMR spectrum (DMSO_d₆) : 4.95 (s, 2H) ; 5.09 (bs, 2H) ; 6.51 (dd, 1H) ; 6.60 (d, 1H) ; 6.68 (s, 1H) ; 6.90 (dd, 1H) ; 6.99-7.05 (m, 2H) ; 7.13 (d, 1H) ; 7.30 (bs, 1H) ; 7.32 (d, 1H) ; 7.90 (bs, 1H), 11.37 (s; 1H).

MS - ESI : 280 [M-H]⁻

10

The starting material was prepared as follows:

**2-Carbamoyl-5-hydroxy-1*H*-indole**

2-Carbamoyl-5-benzylxy-1*H*-indole (1 g ; 3.35 mmol) in solution in methanol (60 ml) was hydrogenated on 10 % Pd/C (0.28 g) for 2 hours. After filtration on celite the solvent was evaporated to give 2-carbamoyl-5-hydroxy-1*H*-indole which was used in the next step without further purification.

Yield : 100 %

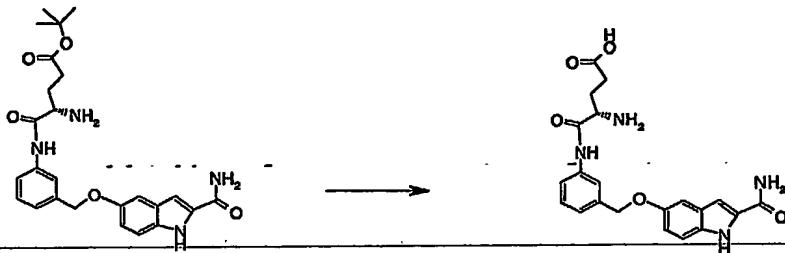
¹H NMR spectrum (DMSO_d₆) : 6.71 (dd, 1H) ; 6.86 (d, 2H) ; 7.20 (d, 1H) ; 7.24 (bs, 1H) ; 7.81 (bs, 1H) ; 8.77 (bs, 1H) ; 11.20 (bs, 1H).

2-Carbamoyl-5-(3-nitrobenzylxy)-1*H*-indole

This was prepared using the same general method as described in Example 1.

Yield : 38 %

²⁵ ¹H NMR spectrum (DMSO_d₆) : 5.27 (d, 2H) ; 6.96 (dd, 1H) ; 7.02 (s, 1H) ; 7.19 (d, 1H) ; 7.30 (bs, 1H) ; 7.34 (d, 1H) ; 7.70 (dd, 1H) ; 7.90 (bs, 1H) ; 7.94 (d, 1H) ; 8.19 (dd, 1H) ; 8.33 (s, 1H) ; 11.42 (bs, 1H).

Example 3**2-Carbamoyl-5-[3-(α -glutamylamino)benzyloxy]-1*H*-indole**

5 2-Carbamoyl-5-[3-(O-*tert*-butyl- α -glutamylamino)benzyloxy]-1*H*-indole (0.29 g ; 0.62 mmol) in solution in CH₂Cl₂ (2 ml) was treated under argon with a HCl (2N) in dioxan (4 ml). The mixture was stirred for one hour, evaporated and the residue was purified on OASIS resin, eluting with H₂O / CH₃CN 80/20 to give 2-carbamoyl-5-[3-(α -glutamylamino)benzyloxy]-1*H*-indole.

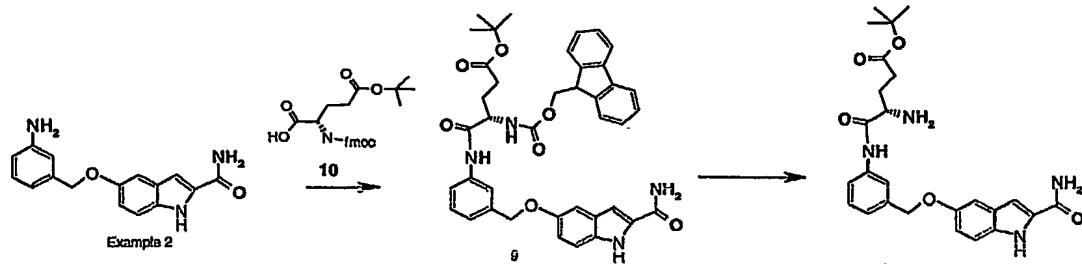
10 Yield : 51 %

¹H NMR spectrum (DMSO-d₆) : 1.97-2.10 (m, 2H) ; 2.34-2.44 (m, 2H) ; 3.94 (t, 1H) ; 5.10 (s, 2H) ; 6.91 (dd, 1H) ; 7.01 (d, 1H) ; 7.16 (d, 1H) ; 7.22 (d, 1H) ; 7.31 (bs, 1H) ; 7.32 (d, 1H) ; 7.37 (dd, 1H) ; 7.60 (d, 1H) ; 7.73 (s, 1H) ; 7.90 (bs, 1H) ; 8.96 (bs, 2H) ; 10.63 (bs, 1H) ; 11.41 (s, 1H).

15 MS - ESI : 409 [M-H]⁻

Elemental analysis	Found	C 54.54	H 5.5	N 12.45
C ₂₁ H ₂₂ N ₄ O ₅ , 0.7 H ₂ O, 1 HCl	Requires	C 54.89	H 5.35	N 12.19

The starting material was prepared as follows:



20

2-Carbamoyl-5-[3-(*tert*-butyl- α -glutamylamino)benzyloxy]-1*H*-indole

A solution of 2-carbamoyl-5-[3-(*tert*-butyl- α -glutamylamino)benzyloxy]-1*H*-indole (Example 3) (0.23 g ; 0.5 mmol) in

1.1 mmol) and **10** (0.467 g ; 1.1 mmol) in CH₂Cl₂ (7 ml). The mixture was stirred overnight under argon atmosphere. After evaporation, the residue was purified by flash chromatography eluting with CH₂Cl₂ and CH₂Cl₂ / Acetone 60/40 to give **9**.

- 5 A solution of **9** (0.51 g ; 0.74 mmol) and piperidine (0.8 ml) in CH₂Cl₂ (10 ml) was stirred at ambient temperature for 3 hours. After evaporation to dryness, the residue was purified by flash chromatography eluting with CH₂Cl₂, CH₂Cl₂ / CH₃CN 90/10 and then with CH₂Cl₂ / CH₃CN / MeOH 45/45/10 to give 2-carbamoyl-5-[3-(*tert*-butyl- α -glutamylamino)benzyloxy]-1*H*-indole (**8**).

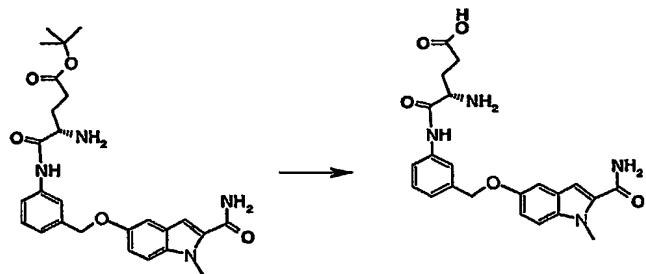
10 Yield : 87 %

¹H NMR spectrum (DMSO-d₆) : 1.38 (s, 9H) ; 1.59-1.76 (m, 1H) ; 1.81-1.93 (m, 1H) ; 2.22-2.36 (m, 2H) ; 3.27-3.34 (m, 2H) ; 5.07 (s, 2H) ; 6.10 (dd, 1H) ; 7.01 (d, 1H) ; 7.12-7.18 (m, 2H) ; 7.26-7.35 (m, 3H) ; 7.59 (d, 1H) ; 7.77 (s, 1H) ; 7.88 (bs, 1H) ; 9.8 (bs, 2H) ; 11.38 (s, 1H).

15

Example 4

1-Methyl-2-carbamoyl-5-[3-(α -glutamylamino)benzyloxy]-1*H*-indole



Example 4

- 20 1-Methyl-2-carbamoyl-5-[3-(α -glutamylamino)benzyloxy]-1*H*-indole was prepared using the same general method as described for Example 3.

Yield : 81 %

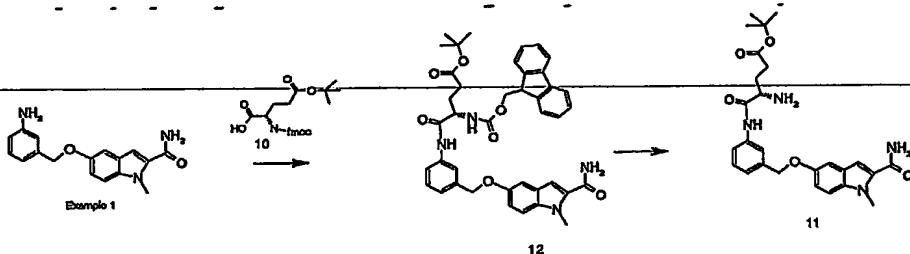
- 25 ¹H NMR spectrum (DMSO-d₆) : 2.02-2.12 (m, 2H) ; 2.36-2.44 (m, 2H) ; 3.97 (s, 3H) ; 4.01 (t, 1H) ; 5.12 (s, 2H) ; 7.00 (dd, 1H) ; 7.03 (s, 1H) ; 7.17 (d, 1H) ; 7.23 (d, 1H) ; 7.31 (bs, 1H) ; 7.38 (dd, 1H) ; 7.44 (d, 1H) ; 7.60 (d, 1H) ; 7.23 (s, 1H) ; 7.90 (bs, 1H) ; 8.83 (bs, 2H) ; 10.72 (s, 1H).

MS - ESI : 423 [M-H]⁺

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Elemental analysis	Found	C 55.04 H 5.53 N 11.49
$\text{C}_{22}\text{H}_{24}\text{N}_4\text{O}_5$, 0.9 H_2O , 1 HCl	Requires	C 55.38 H 5.66 N 11.74

The starting material was prepared using analogous methodology to that described in
 5 Example 3 for the synthesis of 2-carbamoyl-5-[3-(*tert*-butyl- α -glutamylamino)benzyloxy]-
 1*H*-indole.



1-Methyl-2-carbamoyl-5-[3-(*tert*-butyl- α -glutamylamino)benzyloxy]-1*H*-indole.

Yield : 75 % (for the 2 steps)

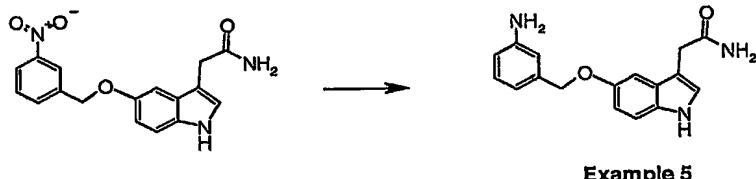
10 ^1H NMR spectrum (DMSO- d_6) : 1.38 (s, 9H) ; 1.64-1.73 (m, 1H) ; 1.83-1.93 (m, 1H) ; 2.22-2.40 (m, 2H) ; 3.34-3.38 (m, 1H) ; 3.96 (s, 3H) ; 5.09 (s, 2H) ; 6.94 -dd, 1H) ; 7.02 (s, 1H) ; 7.12-7.20 (m, 2H) ; 7.25-7.36 (m, 2H) ; 7.43 (d, 1H) ; 7.59 (d, 1H) ; 7.76 (s, 1H) ; 7.89 (bs, 1H) ; 10.0 (bs, 1H), 11.39 (s, 1H).

MS - ESI : 481 [M+H]⁺

15

Example 5

3-Carbamoylmethyl-5-(3-aminobenzyloxy)-1*H*-indole

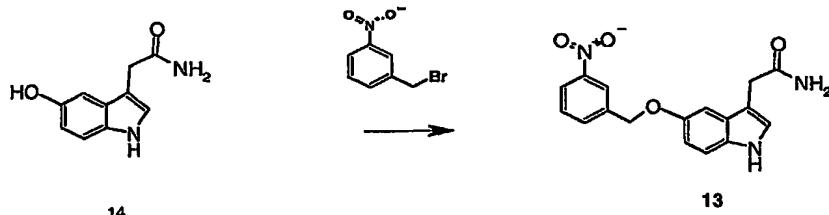


20 3-Carbamoylmethyl-5-(3-aminobenzyloxy)-1*H*-indole was prepared using the same general method as described for Example 1.

Yield : 69 %

^1H NMR spectrum (DMSO- d_6) : 3.43 (s, 2H) ; 4.91 (s, 2H) ; 5.09 (bs, 2H) ; 6.51 (dd, 1H) ; 6.60 (d, 1H) ; 6.70 (s, 1H) ; 6.78 (dd, 1H) ; 6.82 (bs, 1H) ; 7.02 (dd, 1H) ; 7.17 (s, 1H) ; 7.19 (dd, 1H) ; 7.24 (s, 1H) ; 7.25 (bs, 1H) ; 7.27 (dd, 1H) ; 10.71 (s, 1H).

The starting material was prepared using analogous methodology to that described in Example 1 for intermediate 1.



5 3-Carbamoylmethyl-5-(3-nitrobenzyloxy)-1*H*-indole

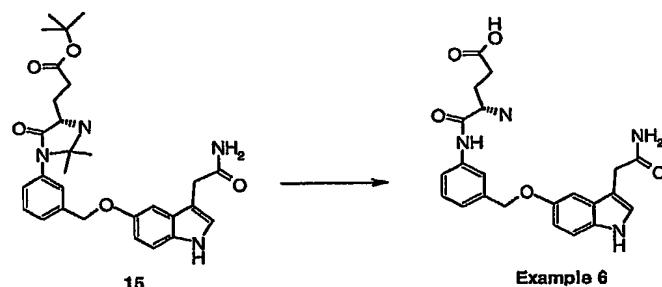
Yield : 58 %

¹H NMR spectrum (DMSO-d₆) : 3.44 (s, 2H) ; 5.26 (d, 2H) ; 6.82 (bs, 1H) ; 6.87 (dd, 1H); 7.18 (d, 1H); 7.20 (dd, 1H) ; 7.28 (bs, 1H) ; 7.28 (d, 1H) ; 7.72 (dd, 1H) ; 7.97 (d, 1H) ; 8.21 (dd, 1H) ; 8.36 (s, 1H) ; 10.77 (s, 1H).

10 MS - ESI : 326 [M+H]⁺

Example 6

3-Carbamoylmethyl-5-(3- α -glutamylbenzyloxy)-1*H*-indole



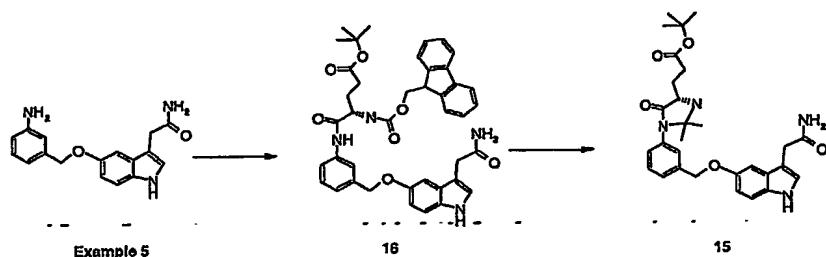
15 3-Carbamoylmethyl-5-(3- α -glutamylbenzyloxy)-1*H*-indole was prepared using the same general method as described in Example 3 by deprotection of 15.

Yield : 27 %

¹H NMR spectrum (DMSO-d₆) : 2.01-2.14 (m, 2H) ; 2.36-2.44 (m, 2H) ; 3.42 (s, 2H) ; 3.98-4.08 (m, 1H) ; 5.08 (s, 2H) ; 6.77-6.83 (m, 2H) ; 7.16 (dd, 1H) ; 7.18 (d, 1H) ; 7.21-7.27 (m, 2H) ; 7.30 (bs, 1H) ; 7.38 (dd, 1H) ; 7.61 (d, 1H) ; 7.74 (s, 1H) ; 8.37 (bs, 2H) ; 10.74 (s, 1H) ; 12.38 (bs, 1H).

MS - ESI : 425 [M+H]⁺

The starting material **15** was prepared using analogous method to that described in Example 3 starting from 3-carbamoylmethyl-5-(3-aminobenzyloxy)-1*H*-indole (Example 5).



Yield : 45.9 %

5 ^1H NMR spectrum (DMSO- d_6) : 1.26 (s, 3H) ; 1.30 (s, 3H) ; 1.42 (s, 9H) ; 1.65-1.78 (m, 1H) ; 1.93-2.05 (m, 1H) ; 2.30-2.45 (m, 2H) ; 3.41 (s, 2H) ; 3.54-3.62 (m, 1H) ; 5.10 (m, 2H) ; 6.79 (bs, 1H) ; 6.81 (dd, 1H) ; 7.10-7.31 (m, 6H) ; 7.46-7.49 (m, 2H) ; 10.71 (s, 1H).

10 Example 7

2-Carbamoyl-6-benzyloxy-1*H*-indole



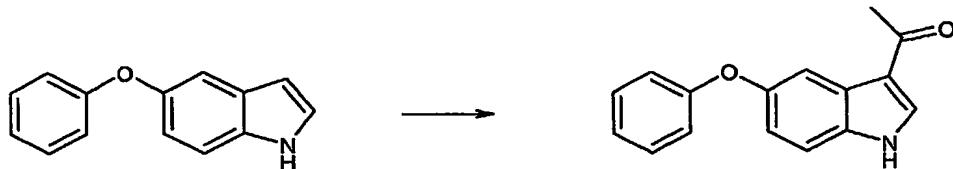
Example 7

A solution of 2-carboxy-6-benzyloxy-1*H*-indole (0.125 g ; 0.468 mmol), oxalyl chloride (0.5 mmol) and DMF (0.2 ml) in CH₂Cl₂ was stirred at room temperature for 2 hours. After evaporation to dryness, the residue was redissolved in acetone ; ammonium acetate (0.074 g ; 0.95 mmol) was added to the solution and the mixture was stirred at room temperature for 1 hour. After evaporation, the residue was purified by flash chromatography eluting with CH₂Cl₂ / acetone 80/20 to give 2-carbamoyl-6-benzyloxy-1*H*-indole.

Yield : 70 %

20 ^1H NMR spectrum (DMSO- d_6) : 5.10 (s, 2H) ; 6.77 (dd, 1H) ; 6.96 (d, 1H) ; 7.04 (d, 1H) ; 7.21 (bs, 1H) ; 7.33 (t, 1H) ; 7.40 (t, 2H) ; 7.46-7.50 (m, 3H) ; 7.82 (bs, 1H) ; 11.34 (s, 1H).

MS - ESI : 265 [M-H]⁻

Example 82-Acetyl-5-phenoxy-1*H*-indole

19

Example 8

EtI (0.23 ml ; 2.87 mmol) was added under argon atmosphere to a suspension of Mg (0.07 g ; 5 2.87 mmol) in ether (3 ml). After stirring for 20 minutes, 19 (0.3 g ; 1.43 mmol) in solution in ether (7 ml) was added. The mixture was stirred for 1 hour, cooled to 10°C and acetyl chloride (0.113 ml ; 1.58 mmol) was added. After stirring at 10°C for 30 minutes, the mixture was poured in ether / H₂O. The organic phase was evaporated and purified by flash chromatography eluting with petroleum ether / AcOEt 60/40 to give 2-acetyl-5-phenoxy-1*H*-indole.

10

Yield : 30 %

¹H NMR spectrum (DMSO-d₆) : 2.41 (s, 3H) ; 6.94 (d, 2H) ; 6.97 (dd, 1H) ; 7.07 (t, 1H) ; 7.35 (dd, 2H) ; 7.49 (d, 2H) ; 7.75 (d, 1H) ; 8.34 (d, 1H) ; 11.99 (bs, 1H).

MS - ESI : 250 [M-H]⁻

15

Example 93-Methoxycarbonyl-5-phenylsulphanyl-1*H*-indole

Example 9

Mg (0.073 g ; 3 mmol) was added under argon atmosphere to a solution of EtI (0.240 ml ; 3 mmol) in ether (4 ml). After stirring for 20 minutes, 20 (0.338 g ; 1.5 mmol) in solution in ether (5 ml) was added. The mixture was cooled to 10°C and methyl chloroformate (0.128 ml ; 1.65 mmol) was added. After stirring for 15 minutes, the mixture was extracted with AcOEt / H₂O. The organic phase was evaporated to give 3-methoxycarbonyl-5-phenylsulphanyl-1*H*-indole.

25 after purification by flash chromatography and elution with petroleum ether / AcOEt 95/5.

Yield : 28 %

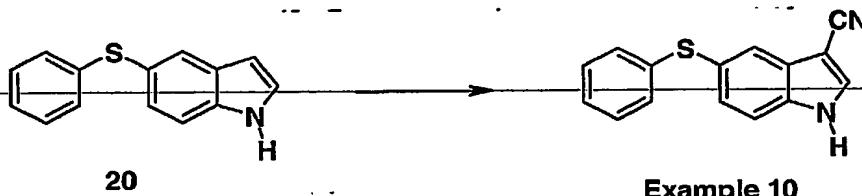
- 52 -

¹H NMR spectrum (DMSO-d₆) : 3.80 (s, 3H) ; 7.12-7.21 (m, 3H) ; 7.26-7.32 (m, 3H) ; 7.56 (d, 1H) ; 8.13-8.18 (m, 1H) ; 12.15 (bs, 1H).

MS - ESI : 282 [M-H]⁻

5 Example 10

3-Cyano-5-phenylsulphanyl-1*H*-indole



20

Example 10

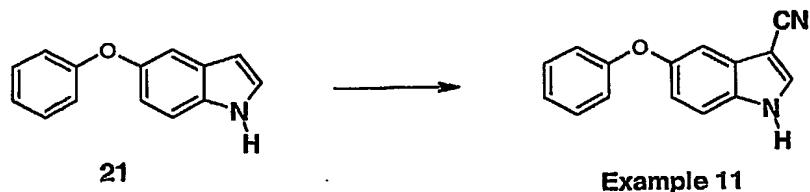
To a stirred solution of 20 (0.45 g ; 2 mmol) in CH₃CN (6 ml) at 0°C was added chlorosulfonyl isocyanate (0.175 ml ; 2 mmol) in solution in CH₃CN (6 ml). Et₃N (0.272 ml ; 10 1.96 mmol) was added over 45 minutes, maintaining the temperature near 0°C. The resulting solution was stirred at room temperature for 2 hours. The solvent was removed and the residue extracted with CH₂Cl₂ / sat. aq. NaHCO₃. The organic phase was purified by flash chromatography eluting with CH₂Cl₂ / EtOH 98/2 to give after trituration in ether 3-cyano-5-phenylsulphanyl-1*H*-indole as a solid.

15 Yield : 38 %

¹H NMR spectrum (CDCl₃) : 7.16-7.23 (m, 1H) ; 7.25-7.29 (m, 4H) ; 7.38-7.46 (m, 2H) ; 7.75 (d, 1H) ; 7.90 (s, 1H) ; 8.75 (bs, 1H).

Example 11

20 **3-Cyano-5-phenoxy-1*H*-indole**



21

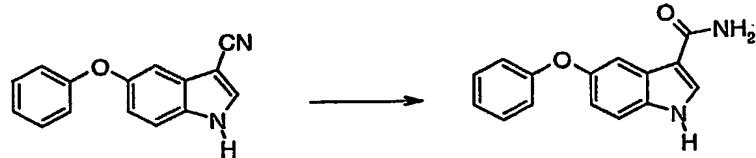
Example 11

3-Cyano-5-phenoxy-1*H*-indole was prepared using the same method as described for Example 10 but starting from 21.

Yield : 62 %

25 ¹H NMR spectrum (DMSO-d₆) : 6.99 (d, 2H) ; 7.04 (d, 1H) ; 7.12 (t, 1H) ; 7.16 (d, 1H) ; 7.38

(d, 2H) ; 7.54 (d, 1H) ; 8.21 (s, 1H) ; 12.12 (bs, 1H).

Example 12**3-Carbamoyl-5-phenoxy-1*H*-indole**

Example 11

Example 12

5

To a solution of 3-Cyano-5-phenoxy-1*H*-indole (Example 11) (0.105 g ; 0.45 mmol) in EtOH (3 ml) and 1N NaOH (1.5 ml) was added H₂O₂ (0.15 ml). The mixture was heated at 50°C for 1 h 30mn, diluted with water, cooled to 5°C and neutralised to pH 7 with 1N HCl.

The mixture was extracted with AcOEt, evaporated to dryness and purified by flash chromatography, eluting with petroleum ether / AcOEt 50/50 to give 3-carbamoyl-5-phenoxy-1*H*-indole.

Yield : 35 %

¹H NMR spectrum (DMSO_d₆) : 6.77 (bs, 1H) ; 6.89-6.96 (m, 3H) ; 7.05 (dd, 1H) ; 7.34 (dd, 2H) ; 7.40 (bs, 1H) ; 7.45 (d, 1H) ; 7.76 (d, 1H) ; 8.06 (d, 1H) ; 11.59 (bs, 1H).

15

MS - ESI : 253 [M-H]⁻

Example 13**3-Carbamoyl-5-(4-hydroxyphenoxy)-1*H*-indole**

20

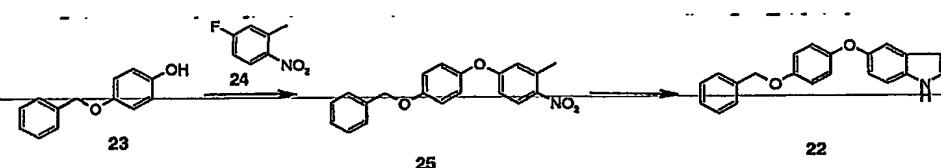
To a stirred solution of 22 (0.25 g ; 0.79 mmol) in ether (10 ml) at 0°C was added chlorosulphonyl isocyanate (0.561 g ; 3.9 mmol). The solution was stirred for 20 minutes. The precipitate was collected by filtration, redissolved in THF (3 ml) and exposed to light for 30 minutes. After evaporation, the residue was taken up in methanol (5 ml). Ammonium acetate (0.498 g ; 0.79 mmol) and Pd/C (0.05 g) were added. The suspension was heated to 80°C for 30 minutes. After filtration and trituration with water, the solid was washed with ether to give 3-carbamoyl-5-(4-hydroxyphenoxy)-1*H*-indole.

Yield : 52 %

¹H NMR (DMSO-d₆) : 6.84 (m, 5H) ; 7.42 (dd, 1H) ; 7.65 (m, 1H) ; 8.08 (s, 1H).

MS - ESI : 267 [M-H]⁻

5 The starting material was prepared as follows :



2-Methyl-4-(4-benzyloxyphenoxy)-nitrobenzene

To a stirred slurry of **23** (25 g ; 124.9 mmol) and K_2CO_3 (34.5 g ; 249.8 mmol) in NMP (100 ml) was added 5-fluoro-2-nitrotoluene (19.56 g ; 126.1 mmol). The mixture was heated to 180°C for 2 hours. The suspension was cooled to room temperature and the product was precipitated into water (300 ml). The solid was filtered and purified by flash chromatography eluting with heptane / CH_2Cl_2 50/50 to give 2-methyl-4-(4-benzyloxyphenoxy)-nitrobenzene (**25**).

15 Yield : 93 %

¹H NMR (DMSO-d₆) : 3.32 (s, 3H) ; 5.11 (s, 2H) ; 6.86 (m, 1H) ; 6.99 (m, 1H) ; 7.11 (m, 5H) ; 7.3-7.5 (m, 5H).

3-Carbamoyl-5-(4-benzyloxypyphenoxy)-1*H*-indole

20 To a stirred solution of **25** (40 g ; 119.4 mmol) and pyrrolidine (8.49 g ; 119.4 mmol) in dry DMF (40 ml) was added N,N-dimethyl formamide dimethyl acetal (42.7 g ; 3.58 mmol). The mixture was heated to 120°C for 2 hours and extracted with ether / H₂O. The organic phase was evaporated to give a red solid. 40 g of the solid in solution in toluene-acetic acid (500 ml ; 3:2) were added to a stirred suspension of Fe (80 g ; 1.43 mol) and silica gel (130 g) in a
 25 mixture of toluene-acetic acid (1.5 l ; 3:2). The reaction mixture was heated to 100°C for 1 hour. The suspension was cooled to ambient temperature and filtered through celite. The filtrate was concentrated to afford an oil which was purified by flash chromatography, eluting with heptane / CH₂Cl₂ 50/50 to give **22**.

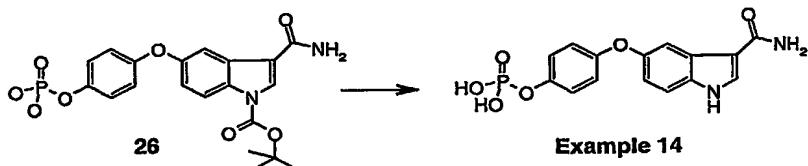
Yield: 43 %

¹H NMR (DMSO-d₆) : 5.05 (s, 2H) ; 6.36 (m, 1H) ; 6.80 (dd, 1H) ; 6.89 (d, 2H) ; 6.97 (d, 2H) ; 7.09 (m, 1H) ; 7.39 (m, 7H).

MS - ESI : 316 [M+H]⁺

5 Example 14

3-Carbamoyl-5-(4-phosphonoxyphenoxy)-1*H*-indole



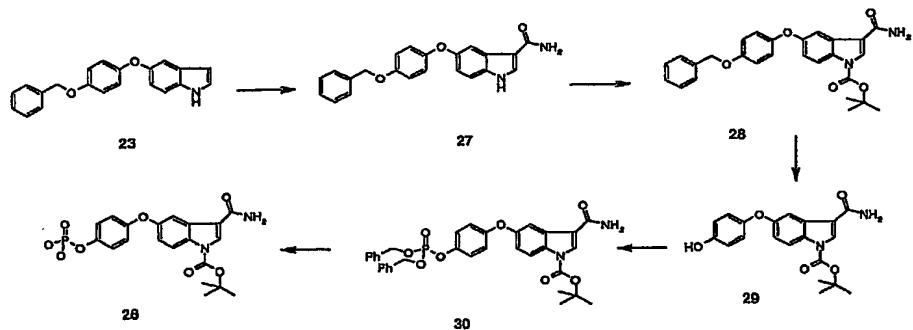
To a mixture of 1-*tert*-butyloxycarbonyl-3-carbamoyl-5-(4-phosphonoxyphenoxy)-1*H*-indole (**26**) (0.47 g ; 1 mmol) in CH₂Cl₂ was added TFA (25 ml). The solution was stirred for 2 hours. The solvent was evaporated and the solid purified on Oasis resin eluting with a 0-30% gradient MeOH/H₂O. After evaporation of methanol, the pH was adjusted to 7.5 with 0.1N NaOH to give after freeze drying 3-carbamoyl-5-(4-phosphonoxyphenoxy)-1*H*-indole.

Yield : 80 %

¹H NMR (DMSO-d₆) : 6.83 (m, 3H) ; 7.16 (d, 2H) ; 7.45 (d, 1H) ; 7.75 (s, 1H) ; 8.09 (s, 1H).

15 MS - ESI : 349 [M+H]⁺

The starting material was prepared as follows:



3-Carbamoyl-5-(4-benzyloxyphenoxy)-1*H*-indole

20 To a stirred solution of **23** (2 g ; 6.3 mmol) in ether (100 ml) at 0°C was added chlorosulphonyl isocyanate (4.49 g ; 32 mmol). The solution was stirred at 0°C for 1 hour. The solid was collected by filtration and dissolved in THF (20 ml). The solution was exposed to light for 1 hour. After evaporation the residue was purified by flash chromatography eluting with CH₂Cl₂ / MeOH 85/15 to give **27**.

25 Yield : 94 %

- 56 -

¹H NMR (DMSO) : 5.08 (s, 2H) ; 6.88 (m, 3H) ; 7.04 (d, 12H) ; 7.41 (m, 5H) ; 7.67 (m, 1H) ; 8.05 (m, 1H).

MS - ESI : 359 [M+H]⁺

5 1-tert-Butyloxycarbonyl-3-carbamoyl-5-(4-benzyloxyphenoxy)-1H-indole

To a stirred solution of **27** (2 g ; 5.6 mmol) in acetonitrile (100 ml) were added DMAP (0.034 g ; 0.28 mmol) and di-*tert*-butyldicarbonate (1.22 g ; 5.6 mmol). The suspension was stirred for 1 hour. The solvent was evaporated and the residue taken up in AcOEt (100 ml), washed with 10 % citric acid and purified by flash chromatography eluting with CH₂Cl₂ / MeOH 95/5

10 to give **28**.

Yield : 61 %

¹H NMR (DMSO) : 1.66 (s, 9H) ; 5.09 (s, 2H) ; 7.05 (m, 5H) ; 7.14 (brs, 1H) ; 7.41 (m, 3H) ; 7.47 (d, 2H) ; 7.75 (m, 1H) ; 7.86 (brs, 1H) ; 8.05 (d, 1H) ; 8.48 (s, 1H).

MS - ESI : 459 [M+H]⁺

15

1-tert-Butyloxycarbonyl-3-carbamoyl-5-(4-hydroxyphenoxy)-1H-indole

To a solution of **28** (1.39 g ; 2.8 mmol) in AcOEt / EtOH (1:2 ; 150 ml), 10 % Pd / C (0.6 g) was added and the suspension was hydrogenated for 2 hours. After filtration on celite the filtrate was evaporated to dryness to give **29**.

20 Yield : 98 %

¹H NMR (DMSO) : 1.64 (s, 9H) ; 6.78 (2H, d) ; 6.88 (d, 2H) ; 7.05 (dd, 1H) ; 7.11 (brs, 1H) ; 7.69 (m, 1H) ; 7.80 (brs, 1H) ; 7.99 (dd, 1H) ; 8.45 (s, 1H) ; 9.30 (brs, 1H).

MS - ESI : 367 [M+H]⁺

25 ¹H NMR (CDCl₃) (DMSO-d₆) : 1.64 (s, 9H) ; 6.78 (d, 2H) ; 6.88 (d, 2H) ; 7.05 (dd, 1H) ; 7.11 (brs, 1H) ; 7.69 (m, 1H) ; 7.80 (brs, 1H) ; 7.99 (dd, 1H) ; 8.45 (s, 1H) ; 9.30 (brs, 1H).

MS - ESI : 369 [M+H]⁺

30 1-tert-Butyloxycarbonyl-3-carbamoyl-5-[4-(di-benzyloxy-phosphonoxy)phenoxy]-1H-indole

To a stirred solution of **29** (1.03 g ; 2.8 mmol), CCl₄ (431 g ; 28 mmol), TPPPA (1.81 g ; 14 mmol) and DMAP (0.28 mmol) in dichloromethane (100 ml) was added triethylamine (1.03 ml)

(4.41 g; 16.8 mmol) over a period of 30 minutes. The mixture was allowed to warm to room temperature and stirred for 1 hour. After evaporation, the residue was taken up in CH_2Cl_2 (200 ml), washed with 10 % citric acid. The organic layer was evaporated and purified by flash chromatography eluting with CH_2Cl_2 / AcOEt 50/50 to give **30**.

5 Yield : 63 %

¹H NMR (DMSO-d₆) : 1.67 (s, 9H) ; 5.19 (s, 4H) ; 7.00 (d, 2H) ; 7.12 (dd, 1H) ; 7.20 (d, 2H) ; 7.39 (9H, m) ; 7.85 (d, 1H) ; 8.10 (d, 2H) ; 8.52 (s, 1H).

MS - ESI : 629 [M+H]⁺

10 1-*tert*-Butyloxycarbonyl-3-carbamoyl-5-(4-phosphonoxypyphenoxy)-1*H*-indole

To a stirred solution of **30** (1.1 g ; 1.8 mmol) in AcOEt / EtOH (30 ml ; 1:1) was added 10 % Pd / C (0.2 g). The suspension was hydrogenated for 1 hour. After filtration of the catalyst the solvent was evaporated to give **26** as a foam.

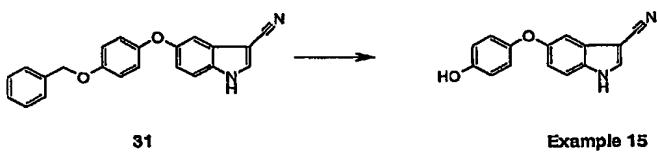
Yield : 58 %

15 ^1H NMR (DMSO- d_6) : 1.65 (s, 9H) ; 6.99 (d, 2H) ; 7.10 (dd, 1H) ; 7.14 (d, 2H) ; 7.91 (m, 1H) ; 7.97 (brs, 1H) ; 8.07 (d, 1H) ; 8.49 (s, 1H).

MS - ESI : 449 [M+H]⁺

Example 15

20 3-Cyano-5-(4-hydroxyphenoxy)-1*H*-indole



To a stirred solution of 31 (0.2 g ; 0.6 mmol) in AcOEt / EtOH (1:1 ; 100 ml) was added 10 % Pd / C (0.05 g). The suspension was hydrogenated for 1 hour. After filtration and evaporation, 25 the residue was purified by flash chromatography eluting with CH₂Cl₂ / MeOH 95/5 to give 3-cyano-5-(4-hydroxyphenoxy)-1*H*-indole.

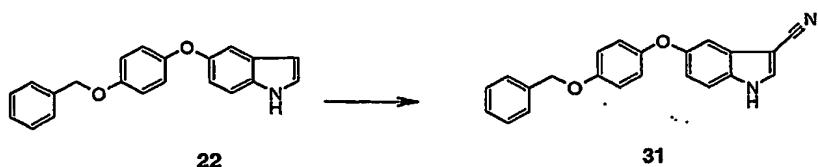
Yield : 100 %

¹H NMR (DMSO-d₆) : 6.78 (m, 3H) ; 6.89 (d, 2H) ; 6.96 (m, 2H) ; 7.53 (dd, 1H) ; 8.22 (s, 1H).

MS - ESI : 249 [M+H]⁺

30

The starting material was prepared as follows:



3-Cyano-5-(4-benzyloxyphenoxy)-1*H*-indole

To a stirred suspension of **22** (0.36 g ; 1.14 mmol) in ether (20 ml) was added chlorosulphonylisocyanate (0.80 g ; 5.7 mmol) at 0°C. The mixture was stirred for 1 hour at 5 0°C. The resulting solid was collected by filtration, redissolved in THF (20 ml) and exposed to light for 1 hour. After evaporation, the residue was purified by flash chromatography eluting with CH₂Cl₂ to give **31** as the by product.

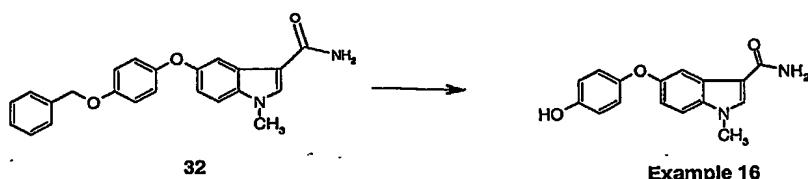
Yield : 17%

¹H NMR (DMSO) 5.06 (s, 2H) ; 6.92 (m, 3H) ; 6.98 (d, 2H) ; 7.39 (m, 6H) ; 7.68 (m, 1H) ; 8.04 (m, 1H)

MS - ESI : 339.53 [M+H]⁺

Example 16

1-Methyl-3-carbamoyl-5-(4-hydroxyphenoxy)-1*H*-indole



15

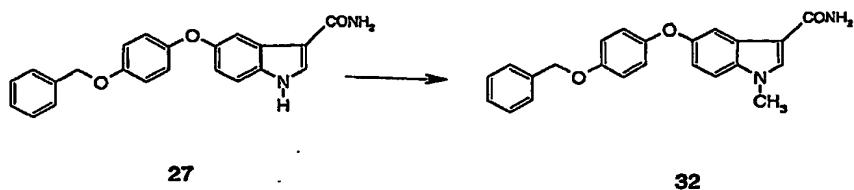
To a stirred solution of **32** (0.2 g ; 0.5 mmol) in EtOH (50 ml) was added Pd (OH)₂ (0.05 g). The suspension was hydrogenated for 1 hour. After filtration and evaporation, the residue was purified by flash chromatography eluting with CH₂Cl₂ / MeOH to give 1-methyl-3-carbamoyl-5-(4-hydroxyphenoxy)-1*H*-indole.

20 Yield : 53 %

¹H NMR (DMSO) : 3.82 (s, 3H) ; 6.74 (d, 2H) ; 6.82 (d, 2H) ; 6.91 (dd, 1H) ; 7.48 (dd, 1H) ; 7.66 (dd, 1H) ; 7.98 (s, 1H) ; 9.21 (s, 1H).

MS - ESI : 281 [M-H]⁻

The starting material was prepared as follows



1-Methyl-3-carbamoyl-5-(4-benzyloxyphenoxy)-1*H*-indole

To a stirred solution of **27** (0.26 g ; 0.76 mmol) in THF (3 ml), at 0°C was added NaH (0.032 g ; 0.8 mmol). MeI (0.108 g ; 0.76 mmol) was added. The reaction was stirred at room temperature for 1 hour. After evaporation the residue was purified by flash chromatography, eluting with CH₂Cl₂ / MeOH 90/10 to give **32**.

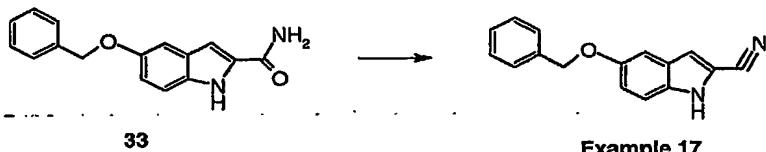
Yield : 70 %

¹H NMR (DMSO) : 3.81 (s, 3H) ; 5.07 (s, 2H) ; 6.91 (m, 3H) ; 6.99 (d, 2H) ; 7.39 (m, 6H) ;
 10 7.68 (m, 1H) ; 7.98 (s, 1H).

MS - ESI : 373 [M+H]⁺

Example 17

2-Cyano-5-benzyloxy-1*H*-indole



15

Trichloromethylchloroformate (0.087 ml ; 0.72 mmol) was added to **33** (0.12 g ; 0.451 mmol) and trimethyl phosphite (0.21 ml). The mixture was heated at 60°C for 5 minutes, diluted with water and extracted with ether. The organic phase was evaporated, the residue was purified by flash chromatography, eluting with CH₂Cl₂ to give 2-cyano-5-benzyloxy-1*H*-indole.

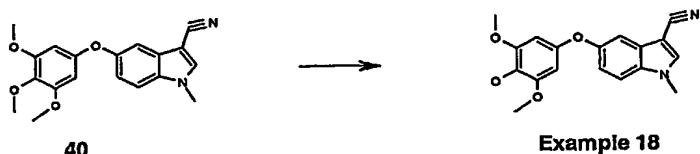
20 Yield : 62 %

¹H NMR spectrum (DMSO-d₆) : 5.11 (s, 2H) ; 7.07 (dd, 1H) ; 7.21 (d, 1H) ; 7.23 (d, 1H) ; 7.33 (dd, 1H) ; 7.36-7.43 (m, 3H) ; 7.47 (d, 2H) ; 12.25 (bs, 1H).

MS - ESI : 247 [M-H]⁻

Example 18

1-Methyl-3-cyano-5-(4-hydroxy-3,5-dimethoxyphenoxy)-1*H*-indole



Example 18

To a solution of 1-methyl-3-cyano-5-(3,4,5-trimethoxyphenoxy)-1*H*-indole (500 mg; 1.47 mmol) in CH₂Cl₂ (10 ml) was added trimethylsilyl iodide (831 μ l ; 5.84 mmol). After stirring at room temperature under argon atmosphere for 2h30, the mixture was poured into cold water, extracted with ethyl acetate. The organic phase was evaporated and the residue was purified by flash chromatography on silica gel, eluting with CH₂Cl₂, Et₂O 1:1 to give 1-methyl-3-cyano-5-(4-hydroxy-3,5-dimethoxyphenoxy)-1*H*-indole: (244 mg)

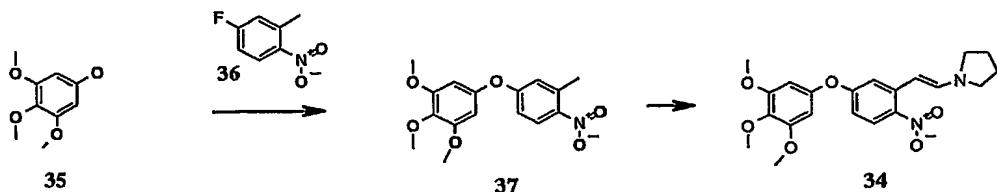
10 Yield: 51%

¹H NMR spectrum (DMSO-d₆): 3.71 (s, 6H); 3.87 (s, 3H); 6.37 (s, 2H); 7.05 (m, 1H); 7.07 (s, 1H); 7.63 (d, 1H); 8.19 (s, 1H); 8.24 (s, 1H).

MS – ESI : 325 [M+H]⁺

Elemental analysis	Found	C 66.19	H 4.98	N 8.47
15 C ₁₈ H ₁₆ N ₂ O ₄	Requires	C 66.66	H 4.97	N 6.64

The starting material was prepared as follows:



2-Methyl-4-(3,4,5-trimethoxyphenoxy)-nitrobenzene

20 A solution of **35** (6g; 32.5 mmol), **36** (5.0 g ; 32.5 mmol) and K_2CO_3 (6.7 g ; 48.5 mmol) in NMP (65 ml) was heated at 80 °C under argon atmosphere for 3h. After cooling, the mixture was poured on cold water and the precipitate was filtrated, washed with water then pentane, and purified by flash chromatography, eluting with CH_2Cl_2 to give **37** as a yellow solid : 9.3g
Yield : 90%.

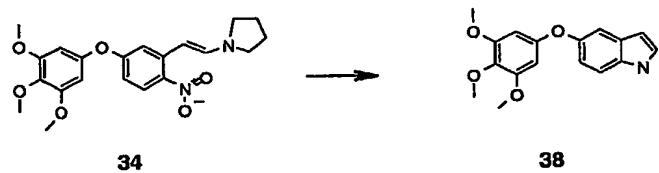
²⁵¹H NMR spectrum (DMSO-d₆) : 2.55 (s, 3H) ; 3.68 (s, 3H) ; 3.72 (s, 6H) ; 6.53 (s, 2H) ; 6.95 (s, 1H).

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To a solution of **37** (9.3 g ; 29.15 mmol) in DMF (15 ml) was added N,N-Dimethylformamide dimethyl acetal (4.5 ml ; 33.8 ml) and pyrrolidine (2.8 ml ; 33.8 mmol). The solution was heated at reflux (110° C) under argon atmosphere for 2h30. After evaporation under vacuum, the residue was dissolved in CH₂Cl₂ (8 ml) and Methanol (6.4 ml). The solution was concentrated to about 6ml and cooled to 5°C. The resulting solid was filtered and washed with cold methanol to give **34** as red crystals : 10.9g

Yield: 94%

¹H NMR spectrum (DMSO-d₆) : 1.89 (m, 4H) ; 3.30 (m, 4H) ; 3.67 (s, 3H) ; 3.72 (s, 6H) ; 5.80 (d, 1H) ; 6.43 (d, 1H) ; 6.47 (s, 2 H) ; 7.27(d, 1 H) ; 7.64 (d, 1H) ; 7.88 (d, 1H).



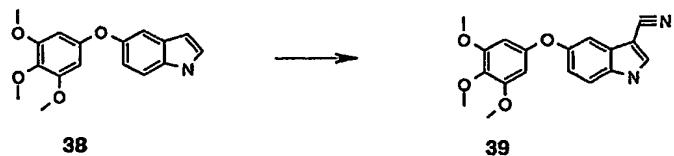
5-(3,4,5-Trimethoxyphenoxy)-1*H*-indole

To a stirred solution of **34** (10.9 g ; 27.25 mmol) in THF (48 ml) and MeOH (48 ml) at 30°C under argon was added excess Raney nickel followed by hydrazine hydrate 85% (2.9 ml). Vigorous gas evolution was observed and temperature rose to 46°C. An additional 2.9 ml of hydrazine hydrate was added 1hr later. The temperature was maintained between 45-50°C for 2h after the last addition. The catalyst was removed by filtration on celite and was washed several times with CH₂Cl₂; after evaporation of the filtrate, the residue was purified by flash chromatography, eluting with AcOEt / Essence G 4:6 then 1:1 to give **38** as a white solid :

3.5 g

Yield : 43%.

¹H NMR spectrum (DMSO-d₆) : 3.63 (s, 3H) ; 3.68 (s, 6H) ; 6.26 (s, 2H) ; 6.41 (d, 1H) ; 6.83 (dd, 1 H) ; 7.19 (d, 1H) ; 7.36 (m, 2H) ; 11.12 (s, 1H).

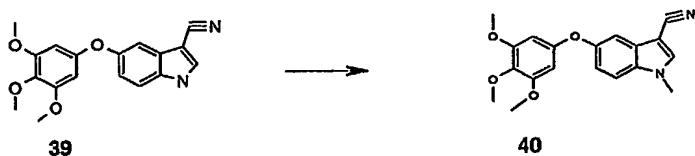


3-Cyano-5-(3,4,5-trimethoxyphenoxy)-1*H*-indole

38 (2.3 g ; 7.69 mmol) in solution in CH₃CN (7ml) and Et₂O (10ml) was treated with Chlorosulfonyl isocyanate (1.33 ml ; 15.38 mmol). After stirring for 1 hour, the solvents were removed and the residue was triturated several times in pentane, dried under vacuum and then taken up in DMF (30ml) ; the solution was stirred at room temperature for 1 hr and poured in cold water. The resulting precipitate was filtrated, dried and purified by flash chromatography on silica gel, eluting with CH₂Cl₂/Et₂O 1:1 to give **39** as a white solid : 2.02 g

Yield: 81%.

10 ^1H NMR spectrum (DMSO- d_6): 3.65 (s, 3H); 3.71 (s, 6H); 6.35 (s, 2H); 7.02 (dd, 1H); 7.18 (d, 1H); 7.56 (d, 1H); 8.26 (s, 1H).

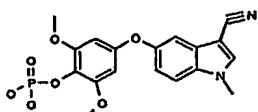
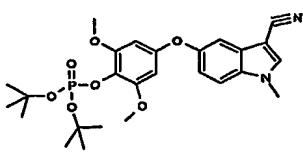


1-Methyl-3-cyano-5-(3,4,5-trimethoxyphenoxy)-1*H*-indole

15 **39** (230mg ; 0.71 mmol) in solution in DMF (0.9 ml) was treated with NaH 60% dispersed in mineral oil (33 mg ; 0.71 mmol). The mixture was stirred under argon atmosphere for 10mn and CH₃I (73μl ; 0.78 mmol) was added . The mixture was stirred under argon atmosphere for 1h and poured into cold water. The resulting white solid was filtrated washed with water and dried under vacuum for a night to give **40** : 230 mg.

20 Yield: 96%

¹H NMR spectrum (DMSO-d₆): 3.66 (s, 3H); 3.71 (s, 6H); 3.89 (s, 3H); 6.35 (s, 2H); 7.10 (dd, 1H); 7.19 (d, 1H); 7.68 (d, 1H); 8.26 (s, 1H).

Example 19**1-Methyl-3-cyano-5-(4-phosphonoxy-3,5-dimethoxyphenoxy)-1*H*-indole**

Example 19

41 (470 mg ; 0.91 mmol) in solution in CH_2Cl_2 (1 ml) was treated under argon with a 2N HCl in dioxan (6.5 ml). The mixture was stirred at room temperature for 2hr, evaporated and the residue was purified by crystallisation in Et_2O / pentane : 250mg

Yield: 68%

^1H NMR spectrum ($\text{DMSO}-\text{d}_6$): 3.68 (s, 6H) ; 3.86 (s, 3H) ; 6.35 (s, 2H) ; 7.10 (dd, 1H) ; 7.20 (s, 1H) ; 7.67 (d, 1H); 8.28 (s, 1H) .

MS – ESI : 405 [M+H] $^+$

Elemental analysis	Found	C 51.00	H 4.45	N 6.50
$\text{C}_{18}\text{H}_{17}\text{N}_2\text{O}_7\text{P}, \text{H}_2\text{O}$	Requires	C 51.19	H 4.53	N 6.63

The starting material 41 was prepared as follows:



Example 18

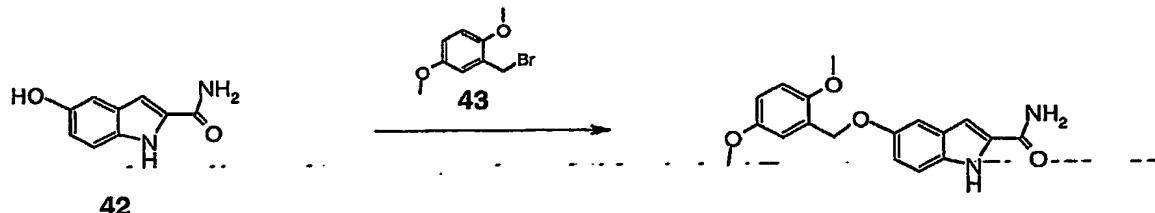
41

1-Methyl-3-cyano-5-[4-(di-*tert*-butylphosphonoxy)-3,5-dimethoxyphenoxy]-1*H*-indole

To a solution of M584994 (870 mg ; 2.68 mmol) in THF (4ml) was added tetrazole (376 mg ; 5.37 mmol) and di-*tert*-butyl-N,N-diethylphosphoramidate (900 μ l ; 3.22 mmol). The mixture was stirred at room temperature for 2hr, cooled to -40°C and a solution of MCPBA (659mg ; 2.95 mmol) in CH_2Cl_2 (5ml) was added . After stirring at 0°C for 1h, the mixture was poured on cold water, washed with NaHSO_3 and extracted with CH_2Cl_2 . The organic phase was washed with cold 2N NaOH and evaporated. The residue was purified by flash chromatography on silica gel, eluting with CH_2Cl_2 , AcOEt 8:2 then with CH_2Cl_2 , AcOEt / Acetone 8 : 1.5 : 0.5 to give 5 : 471 mg

Yield: 33,6%

^1H NMR spectrum ($\text{DMSO}-\text{d}_6$): 1.45 (s, 18 H) ; 3.70 (s, 6H) ; 3.89 (s, 3H) ; 6.39 (s, 2H) ; 7.11 (dd, 1H) ; 7.18 (s, 1 H) ; 7.70 (d, 1H); 8.27 (s, 1H)

Example 20**2-Carbamoyl-5-(2,5-dimethoxybenzyloxy)-1*H*-indole**Example 20

5

A mixture of 42 (0.265 g ; 1.5 mmol), 43 (0.381 g ; 1.65 mmol) and Cs₂CO₃ (0.536 g ; 1.65 mmol) in acetonitrile (20 ml) was heated at 90°C under argon atmosphere for 2 hours. After evaporation to dryness the residue was purified by flash chromatography eluting with a 0-30 % gradient of acetone / CH₂Cl₂ to give the title compound.

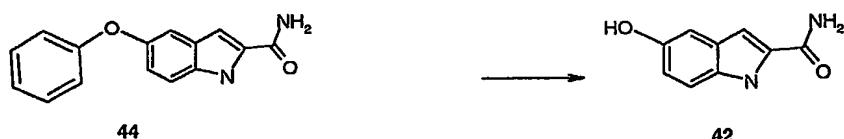
10 Yield : 24 %

¹H NMR Spectrum (DMSO-d₆) : 3.69 (s, 3H) ; 3.74 (s, 3H) ; 5.02 (s, 2H) ; 6.8-6.93 (m, 3H) ; 6.95-7.08 (m, 2H) ; 7.12 (d, 1H) ; 7.28 (s, 1H) ; 7.30 (d, 1H) ; 7.88 (s, 1H) ; 11.36 (s, 1H).

MS-ESI : 327 [M+H]⁺

15

The starting material was prepared as follows :

2-Carbamoyl-5-hydroxy-1*H*-indole

A solution of 44 (1 g ; 3.35 mmol) in Methanol (60 ml) was hydrogenated under medium pressure (45 psi) over 10 % palladium on carbon (0.5 g) for 2 hours. After filtration of the catalyst on celite, the filtrate was evaporated to dryness to give 42 which was used without further purification.

Yield : 95 %

¹H NMR Spectrum (DMSO-d₆) : 6.71-6.74 (m, 1H) ; 6.8-6.95 (m, 2H) ; 7.1-7.3 (m, 2H) ; 7.83 (broad t, 1H) ; 8.73 (broad t, 1H) ; 11.21 (s, 1H).

25

Example 21**3-Cyano-5-(3,4-dimethoxyphenylsulphanyl)-1*H*-indole**

A solution of chlorosulfonyl isocyanate (0.365 ml ; 4.2 mmol) in acetonitrile (1.5 ml) was
 5 added at 0°C to a solution of 45 (1.2 g ; 4.2 mmol) in acetonitrile (12 ml). The mixture was
 stirred under argon atmosphere for 1 hour at 0°C. A solution of triethylamine (0.571 ml) in
 acetonitrile (3 ml) was then added at 0°C ; the mixture was allowed to warm up and stirred at
 room temperature for 3 hours. After evaporation the residue was taken up in CH₂Cl₂. The
 organic phase was washed with diluted NaHCO₃, evaporated and purified by flash
 10 chromatography eluting with CH₂Cl₂ to give the title compound.

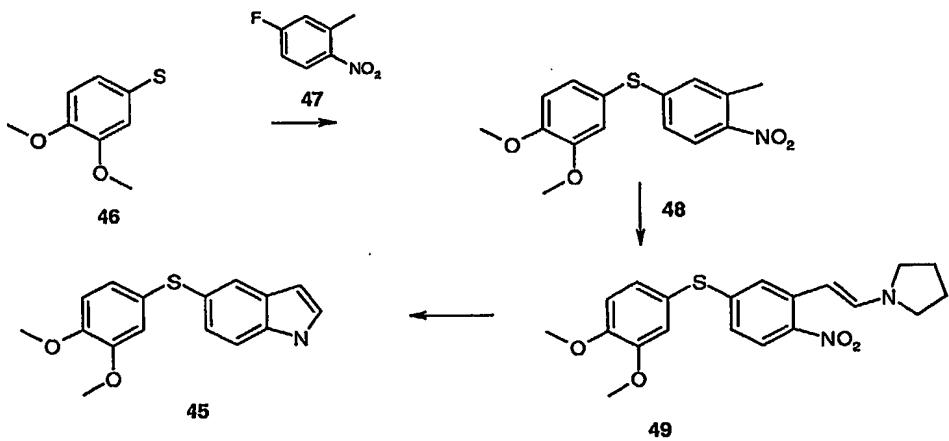
Yield : 52 %

MS-ESI : 311 [M+H]⁺

¹H NMR Spectrum (DMSO-d₆) : 3.72 (s, 3H) ; 3.77 (s, 3H) ; 6.93-7.023 (m, 3H) ; 7.22 (d, 1H)
 ; 7.53 (m, 2H) ; 8.28 (s, 1H) ; 12.0 (s, 1H).

15

The starting material was prepared as follows :

2-Methyl-4-(3,4-dimethoxyphenylsulphanyl)-nitrobenzene

To a solution of 46 (3.4 g ; 20 mmol) and 47 (3.10 g ; 30 mmol) in NMP (40 ml) was added
 20 K₂CO₃ (4.14 g ; 30 mmol). The mixture was stirred at 80°C for 2 hours under argon
 atmosphere. After extraction with a mixture of Ether / Ethylacetate, the organic phase was

evaporated and purified by flash chromatography, eluting with petroleum ether and petroleum ether / ethylacetate 7 : 3 to give **48**.

Yield : 83 %

¹H NMR Spectrum (DMSO_d₆) : 2.48 (s, 3H) ; 3.74 (s, 3H) ; 3.77 (s, 3H) ; 6.97 (dd, 1H) ; 7.10-
5 7.20 (m, 4H) ; 7.9 (d, 1H).

2-(pyrrolidine-1-ylpropanyl-2-ylidene-4-(3,4-dimethoxyphenylsulphanyl)-nitrobenzene

To a solution of **48** (5.1 g ; 16.7 mmol) in DMF (9 ml) was added N,N-dimethylformamide dimethylacetal (2.57 ml ; 19.3 mmol) and pyrrolidine (1.61 ml ; 19.3 mmol). The mixture was 10 heated at reflux under argon atmosphere for 3 hours. After extraction with AcOEt, the organic phase was evaporated to give **49** which was used in the next step without purification.

Yield : 92 %.

5-(3,4-Dimethoxyphenylsulphanyl)-1*H*-indole

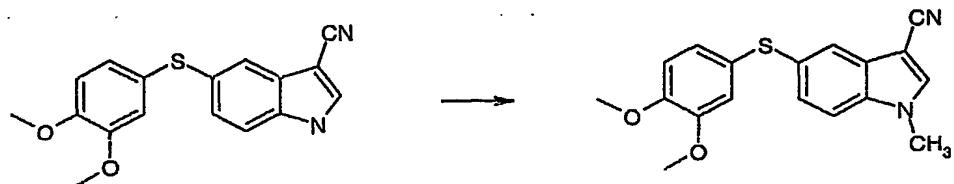
15 To a solution of **49** (5.9 g ; 15.2 mmol) in MeOH / THF (30 ml / 30 ml) was added Raney Nickel (3 g). The mixture was heated at 48°C, under argon atmosphere and a solution of hydrazine hydrate (1.52 g ; 30.4 mmol) in THF / MeOH 30 ml / 30 ml was added dropwise. After heating at 48°C for 2 hours, the mixture was filtered on celite. The filtrate was evaporated and purified by flash chromatography eluting with AcOEt / petroleum ether 30/70 20 to give **45**.

Yield : 35 %

¹H NMR Spectrum : 3.69 (s, 3H) ; 3.73 (s, 3H) ; 6.43 (s, 1H) ; 6.77 (dd, 1H) ; 6.90-6.92 (m, 2H) ; 7.11 (dd, 1H) ; 7.39 (m, 2H) ; 7.64 (s, 1H) ; 11.25 (s, 1H).

25 Example 22

1-Methyl-3-cyano-5-(3,4-dimethoxyphenylsulphanyl)-1*H*-indole



Example 21

Example 22

To a solution of Example 21 (0.2 g ; 0.64 mmol) in DMF (0.8 ml) at 0°C was added NaH (0.03 g ; 0.64 mmol) and CH₃I (0.06 ml ; 0.64 mmol). The mixture was stirred at ambient temperature for 2 hours. After extraction with AcOEt the organic phase was evaporated and purified by flash chromatography eluting with petroleum ether / AcOEt 60/40 to give the title compound.

Yield : 77 %

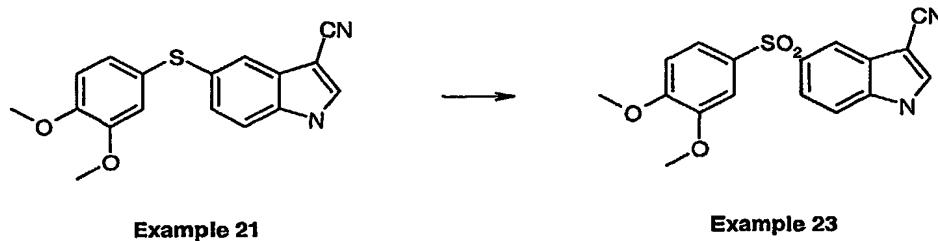
¹H NMR Spectrum (DMSO_d₆) : 3.72 (s, 3H) ; 3.77 (s, 3H) ; 3.87 (s, 3H) ; 6.98-7.03 (m, 3H) ; 7.30 (dd, 1H) ; 7.52 (s, 1H) ; 7.63 (d, 1H) ; 8.28 (s, 1H).

MS-ESI : 325 [M+H]⁺

10

Example 23

3-Cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole

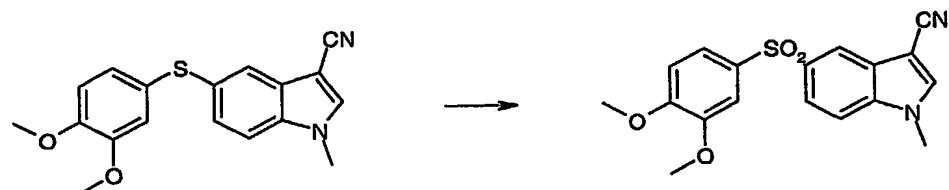


15 To a solution of Example 21 (0.13 g ; 0.42 mmol) in CH₂Cl₂ (10 ml) and CH₃CN (2 ml) was added a solution of MCPBA (0.19 g ; 0.88 mmol) in chloroform (5 ml). The mixture was heated at ambient temperature for one hour. After evaporation to dryness, the solid was triturated in ether and dried to give the title compound.

Yield : 59 %

20 ¹H NMR (DMSO_d₆) : 3.82 (s, 3H) ; 3.84 (s, 3H) ; 7.15 (d, 1H) ; 7.45 (d, 1H) ; 7.61 (dd, 1H) ; 7.73 (d, 1H) ; 7.81 (dd, 1H) ; 8.25 (s, 1H) ; 8.48 (s, 1H).

MS-ESI : 341 [M-H]⁻

Example 24**1-Methyl-3-cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole**

Example 22

Example 24

5 The compound was prepared following the method described for example 23 but starting with compound of example 22.

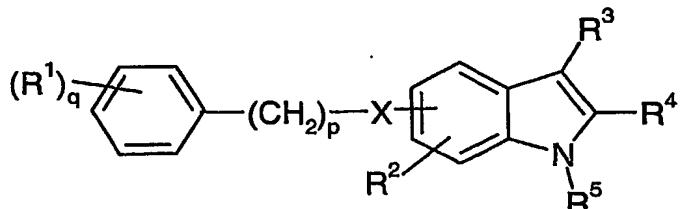
Yield : 77 %.

¹H NMR (DMSO_d₆) : 3.78 (s, 3H) ; 3.82 (s, 3H) ; 3.91 (s, 3H) ; 7.13 (d, 1H) ; 7.46 (s, 1H) ; 7.60 (dd, 1H) ; 7.84-7.91 (m, 2H) ; 8.25 (s, 1H) ; 8.48 (s, 1H).

10 MS-ESI : 357 [M+H]⁺

Claims

1. The use of a compound of Formula (I), for the manufacture of a medicament to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis,



5

Formula (I)

wherein:

R¹ is independently selected from halo, hydroxy, amino, alkanoylamino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;

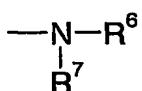
10

X is selected from: —O—, —S—, —SO— or —SO₂—;

R² is selected from: hydrogen, C₁₋₄alkyl or C₁₋₄alkoxy;

15

R³ and R⁴ are independently selected from: hydrogen, C₁₋₄alkyl, C₁₋₄alkanoyl, C₁₋₄alkoxycarbonyl, C₁₋₄alkoxycarbonylC₁₋₄alkyl, C₁₋₄alkoxycarbonylamino, amino, aminoC₁₋₄alkyl, carbamoyl, carbamoylC₁₋₄alkyl, cyano, cyanoC₁₋₄alkyl, hydroxy, hydroxyC₁₋₄alkyl, or a group of Formula (II):



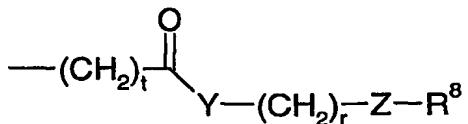
Formula (II)

R⁶ is hydrogen or C₁₋₄alkyl;

20

R⁵ and R⁷ are independently selected from hydrogen, C₁₋₄alkyl or a group of

Formula (III):



Formula (III)

wherein Y is selected from —NH—, —O— or a bond;

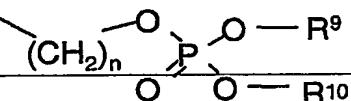
25

Z is selected from —NH—, —O—, —C(O)— or a bond;

r is an integer from 0 to 4;

t is an integer from 0 to 1;

R⁸ is hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy, aryl, 5 or 6 membered heterocyclyl, 5- or 6-membered heteroaryl, wherein aryl, heteroaryl or heterocyclyl are optionally substituted by C₁₋₄alkyl, C₁₋₄alkoxy, or a group of Formula (IV):



Formula (IV)

wherein n is an integer from 1 to 6, and;

R⁹ and R¹⁰ are independently selected from hydrogen,

C₁₋₄alkyl or aryl;

p is an integer between 0 and 1; and

q is an integer between 0 and 3;

with the proviso that:

(i) when R³ is cyano then R⁴ cannot be a group of Formula (II), and

(ii) when q is 0, R³ is cyano and X is —S— then R⁴ is other than amino; or a salt, pro-drug or solvate thereof.

2. The use of a compound of Formula (I), or a salt, pro-drug or solvate thereof, for the manufacture of a medicament to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis, according to

Claim 1 wherein: R¹ is hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified.

3. The use of a compound of Formula (I), or a salt, pro-drug or solvate thereof, for the manufacture of a medicament to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis, according to either Claim 1 or Claim 2 wherein: X is —O— or —S—.

4. The use of a compound of Formula (I), or a salt, pro-drug or solvate thereof, for the

angiogenesis and/or any disease state associated with angiogenesis, according to any one of the preceding claims wherein: R³ is cyano.

5. The use of a compound of Formula (I), or a salt, pro-drug or solvate thereof, for the manufacture of a medicament to inhibit and/or reverse and/or alleviate symptoms of angiogenesis and/or any disease state associated with angiogenesis, according to
5 Claim 1 wherein:

R¹ is selected from hydroxy, amino, —OPO₃H₂, or C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue;
R² is hydrogen;

10 X is selected from: —O—, —S—, —SO— or —SO₂—;

p is 0 or 1;

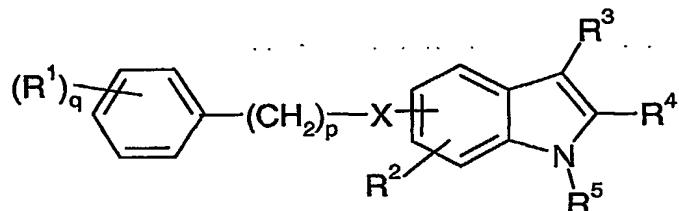
q is an integer between 1 and 3;

R³ is selected from: hydrogen, cyano, carbamoyl, carbamoylC₁₋₄alkyl, C₁₋₄alkanoy, or C₁₋₄alkoxycarbonyl;

15 R⁴ is selected from: hydrogen, cyano or carbamoyl; and

R⁵ is hydrogen or C₁₋₄alkyl.

6. The use of a compound of Formula (V) as a medicament,



Formula (V)

20 wherein:

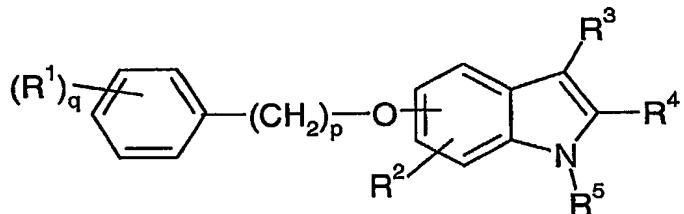
R¹, R², R³, R⁴, R⁵, X, p and q are as defined in Claim 1,

with the proviso that:

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II);
- (ii) when q is 0 and X is —S— then R³ is other than cyano;
- 25 (iii) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is cyanomethyl and R⁴ is hydrogen or R³ is hydrogen and R⁴ is cyanomethyl then R⁵ cannot be hydrogen methyl or acetyl;

- (iv) when $(R^1)_q$ is 4-methoxy, 4-amino, or 3,4,5-trimethoxy, p is 0 or 1, R^2 is hydrogen, 4-chloro or 5-methoxy, R^3 is hydrogen, cyanomethyl or 2-aminoethyl, R^4 is hydrogen or ethoxycarbonyl then R^5 cannot be hydrogen or methyl;
- 5 (v) when q is 0, p is 0 or 1, R^2 is hydrogen, 7-methyl, 5-methoxy or 6-methoxy, R^3 is aminomethyl, 2-aminoethyl, 2-aminopropyl or 2-aminobutyl, R^4 is hydrogen or methyl, then R^5 cannot be hydrogen, methyl, n-butyl or acetyl;
- (vi) when q is 0, p is 0 or 1, R^2 is hydrogen, R^3 is methyl, ethyl, hydroxy, hydroxymethyl, 2-hydroxyethyl or 1-methylethoxy, R^4 is hydrogen, methyl, ethoxycarbonyl or *tert*-butoxycarbonyl or carbamoyl, then R^5 cannot be hydrogen, methyl or acetyl;
- 10 (vii) when q is 0, p is 0 or 1, R^2 is hydrogen or 5-methoxy, R^3 is hydrogen or bromo, R^4 is hydrogen, methyl, methoxycarbonyl, ethoxycarbonyl, *tert*-butoxycarbonyl or hydroxymethyl then R^5 cannot be methyl or n-butyl; and
- 15 (viii) when q is 0, p is 1, R^2 is hydrogen, 6-methyl, 7-methyl, or 5-methoxy, R^3 is hydrogen, R^4 is hydrogen, methoxycarbonyl, ethoxycarbonyl, or 2-aminoethyl, then R^5 cannot be hydrogen; or a salt, pro-drug or solvate thereof.

20 7. A compound of Formula (VI),



Formula (VI)

wherein:

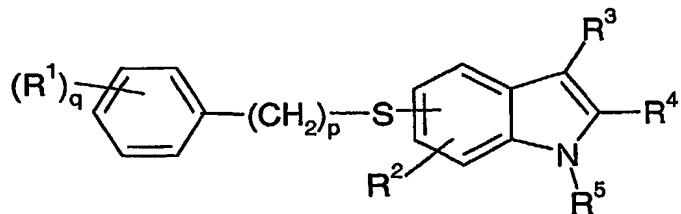
p, q, R^1 , R^2 , R^3 , R^4 , and R^5 are as defined in Claim 1;

25 with the proviso that

- (i) when R^3 is cyano then R^4 cannot be a group of Formula (II);

- (ii) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is cyanomethyl and R⁴ is hydrogen or R³ is hydrogen and R⁴ is cyanomethyl then R⁵ cannot be hydrogen methyl or acetyl;
- 5 (iii) when (R¹)_q is 4-methoxy, 4-amino, or 3,4,5-trimethoxy, p is 0 or 1, R² is hydrogen, 4-chloro or 5-methoxy, R³ is hydrogen, cyanomethyl or 2-aminoethyl, R⁴ is hydrogen or ethoxycarbonyl then R⁵ cannot be hydrogen or methyl;
- 10 (iv) when q is 0, p is 0 or 1, R² is hydrogen, 7-methyl, 5-methoxy or 6-methoxy, R³ is aminomethyl, 2-aminoethyl, 2-aminopropyl or 2-aminobutyl, R⁴ is hydrogen or methyl, then R⁵ cannot be hydrogen, methyl, n-butyl or acetyl;
- 15 (v) when q is 0, p is 1, R² is hydrogen, R³ is methyl, ethyl, hydroxy, hydroxymethyl, 2-hydroxyethyl or 1-methylethoxy, R⁴ is hydrogen, methyl, ethoxycarbonyl or *tert*-butoxycarbonyl or carbamoyl, then R⁵ cannot be hydrogen, methyl or acetyl;
- (vi) when q is 0, p is 1, R² is hydrogen or 5-methoxy, R³ is hydrogen or bromo, R⁴ is hydrogen, methyl, methoxycarbonyl, ethoxycarbonyl, *tert*-butoxycarbonyl or hydroxymethyl then R⁵ cannot be methyl or n-butyl; and
- 20 (vii) when q is 0, p is 1, R² is hydrogen, 6-methyl, 7-methyl, or 5-methoxy, R³ is hydrogen, R⁴ is methoxycarbonyl, ethoxycarbonyl, or 2-aminoethyl, then R⁵ cannot be hydrogen;
- or a salt, pro-drug or solvate thereof.

8. A compound of Formula (VII),



Formula (VII)

25 wherein:

p, q, R¹, R², R³, R⁴, and R⁵ are as defined in Claim 1;

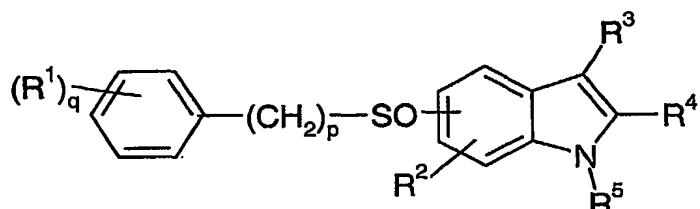
with the proviso that

- (i) when R³ is cyano then R⁴ cannot be a group of Formula (II);

- (ii) when q is 0 or $(R^1)_q$ is 4-amino, p is 0 or 1, R^2 is hydrogen, R^3 is hydrogen, R^4 is hydrogen or ethoxycarbonyl, then R^5 cannot be hydrogen;
- (iii) when q is 0, p is 0 or 1, R^2 is hydrogen, R^3 is hydrogen, methyl, cyano, 2-aminoethyl, 1-methyl-2-aminoethyl or 2-aminopropyl, R^4 is hydrogen or amino, then R^5 cannot be hydrogen; and
- (iv) when q is 0, p is 0, R^2 is hydrogen, R^4 is ethoxycarbonyl, then R^5 cannot be hydrogen;

or a salt, pro-drug or solvate thereof.

9. A compound of Formula (VIIa),



10

Formula (VIIa)

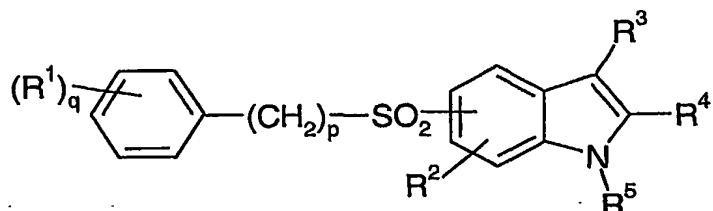
wherein:

p, q, R^1 , R^2 , R^3 , R^4 , and R^5 are as defined in Claim 1;

with the proviso that

- 15 (i) when R^3 is cyano then R^4 cannot be a group of Formula (II);
or a salt, pro-drug or solvate thereof.

10. A compound of Formula (VIIb),



20

Formula (VIIb)

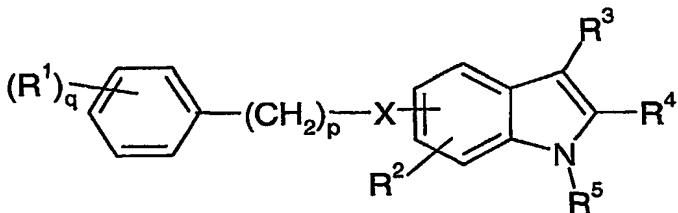
wherein:

p, q, R^1 , R^2 , R^3 , R^4 , and R^5 are as defined in Claim 1;

with the proviso that

- (ii) when $(R^1)_q$ is 4-methyl or 4-chloro, p is 0, R² is hydrogen, R³ is hydrogen, R⁴ is hydrogen or ethoxycarbonyl, then R⁵ cannot be hydrogen; or a salt, pro-drug or solvate thereof.

5 11. A compound of Formula (VIId),



Formula (VIId)

wherein:

- R¹ is independently selected from hydroxy, amino, alkanoylamino, —OPO₃H₂, or
10 C₁₋₄alkoxy, wherein the amino group is optionally substituted by an amino acid residue and the hydroxy group is optionally esterified;
- X, p, R², R³, R⁴, and R⁵ are as defined in Claim 1;
- q is an integer between 1 and 3;
- with the proviso that
- 15 (i) when R³ is cyano then R⁴ cannot be a group of Formula (II); and
- (ii) when $(R^1)_q$ is 4-methoxy, 4-amino, 4-chloro or 3,4,5-trimethoxy, p is 0 or 1, R² is hydrogen, 4-chloro or 5-methoxy, R³ is hydrogen, cyanomethyl or 2-aminoethyl, R⁴ is hydrogen or ethoxycarbonyl, then R⁵ cannot be hydrogen or methyl;
- 20 or a salt, pro-drug or solvate thereof.

12. A compound selected from:

3-cyano-5-phenylsulphanyl-1*H*-indole;

3-cyano-5-phenoxy-1*H*-indole;

25 3-cyano-5-(4-hydroxyphenoxy)-1*H*-indole; and

2-cyano-5-benzyloxy-1*H*-indole;

1-methyl-3-cyano-5-(4-hydroxy-3,5-dimethoxyphenoxy)-1*H*-indole;

1-methyl-3-cyano-5-(4-phosphonoxy-3,5-dimethoxyphenoxy)-1*H*-indole;

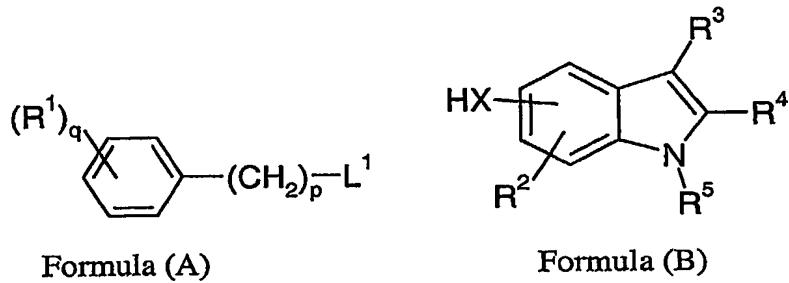
3-cyano-5-(3,4-dimethoxyphenylsulphanyl)-1*H*-indole;

1-methyl-3-cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole;
 3-cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole; and
 1-methyl-3-cyano-5-(3,4-dimethoxyphenylsulphonyl)-1*H*-indole;
 or salt, pro-drug or solvate thereof.

5 13. A pharmaceutical composition comprising a compound according to any one of Claims 6 to 12 or a pharmaceutically acceptable salt, solvate or pro-drug thereof.

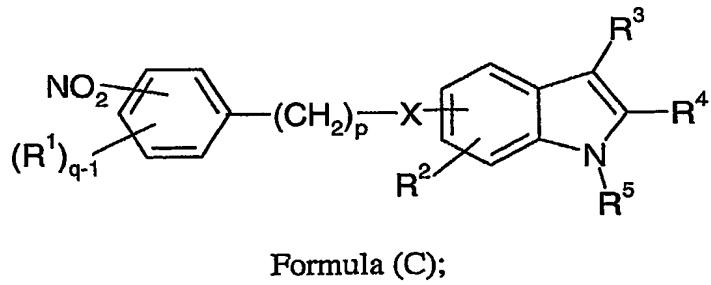
14. A process for preparing a compound of Formula (I), or salt, solvate or pro-drug thereof, which process (wherein R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, X, Y, Z, n, p, q, r and t are unless otherwise specified as defined in Claim 1, comprising:

10 a) for compounds of Formula (I) wherein X is —O—, or —S—, reacting a compound of Formula (A) with a compound of Formula (B),



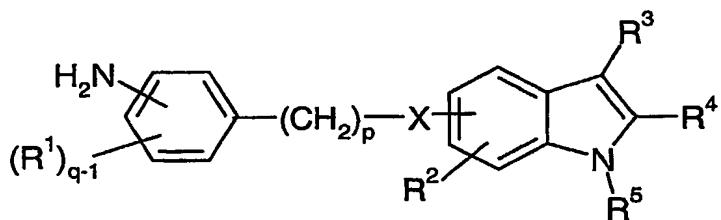
wherein L¹ is a leaving group;

15 b) for compounds of Formula (I) in which R¹ is amino, reduction of a compound of Formula (C):



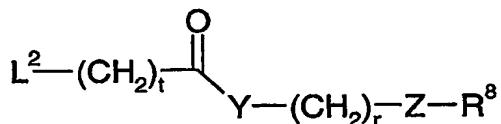
Formula (C);

- c) for compounds of Formula (I) wherein R⁵ is C₁₋₄alkyl, reacting a compound of Formula (I) wherein R⁵ is hydrogen with a suitable alkylhalide;
- d) for compounds of Formula (I) wherein R¹ is an amino group substituted by an amino acid residue, reacting a compound of Formula (D) with an amino acid,



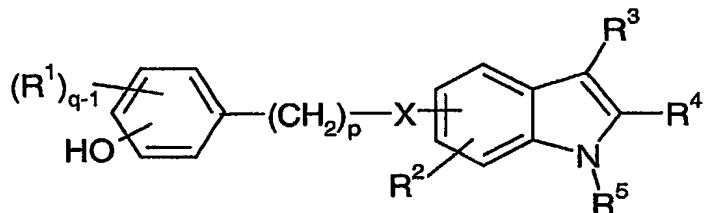
Formula (D);

- 5 e) for compounds of Formula (I) in which R⁵ is a group of Formula (II) and R⁷ is a group of Formula (III), reacting a compounds of Formula (I) in which R⁵ is a group of Formula (II) and R⁷ is hydrogen with compounds of Formula (E) below, in which L² is a leaving group:



Formula (E);

- 10 f) for compounds of Formula (I) in which R⁴ is hydrogen, reacting a compounds of Formula (I) in which R³ is hydrogen and R⁴ is hydrogen with a compounds of L³R³ in which L³ is a leaving group; and
- g) for compounds of Formula (I) in which R¹ is an esterified hydroxyl group, reacting a compound of Formula (F) with an appropriate carboxylic acid or carboxylic acid derivative;



Formula (F)

15

and thereafter if necessary:

- 20 i) converting a compound of Formula (I) into another compound of Formula (I);
ii) removing any protecting groups;
iii) forming a salt, pro-drug or solvate.

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